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**(54) Title:** RECOMBINANT HEPATITIS C VIRUS RNA REPLICASE**(57) Abstract**

A recombinant RNA-dependent RNA polymerase of hepatitis C virus (r-HCV-RDRP) coding DNA was cloned and expressed yielding active enzyme *in vitro*. The r-HCV-RDRP can include up to 20 added amino acids and up to nine deleted or substituted amino acids at the NH<sub>2</sub>-terminus of the encoded amino acid sequence. The invention provides method to solubilize r-HCV-RDRP from a host cell lysate and purified r-HCV-RDRP. Methods for screening for inhibitors of r-HCV-RDRP *in vitro*, for making stably transfected mammalian cells expressing r-HCV-RDRP and for *in vivo* testing of r-HCV-RDRP inhibitors *in vivo* are disclosed. The invention provides antibodies to r-HCV-RDRP and methods for detecting antibodies to HCV-RDRP in serum of human patients.

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## RECOMBINANT HEPATITIS C VIRUS RNA REPLICASE

This application claims priority from U.S. Provisional Application Serial No. 60/004,383, filed September 27, 1995.

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Field of the Invention

The present invention relates Hepatitis-C virus (HCV), specifically to expression and purification of an RNA-dependent RNA polymerase (RDRP) encoded by the HCV genome, to antibodies directed against HCV-RDRP and to methods of using the enzyme to diagnose chronic HCV infections and to screen for antiviral agents effective against HCV.

Background of the Invention

HCV is the major causative agent for post-transfusion and for sporadic non A, non B hepatitis (Alter, H.J. (1990) *J. Gastro. Hepatol.* **1**:78-94; Dienstag, J.L. (1983) *Gastro* **85**:439-462). Despite improved screening, HCV still accounts for at least 25% of the acute viral hepatitis in many countries (Alter, H.J. (1990) *supra*; Dienstag, J.L. (1983) *supra*; Alter, M.J. et al. (1990a) *J.A.M.A.* **264**:2231-2235; Alter, M.J. et al (1990b) *N. Engl. J. Med.* **327**:1899-1905; Alter, M.J. et al .(1990b) *N. Engl. J. Med.* **321**:1494-1500). Infection by HCV is insidious in a high proportion of chronically infected (and infectious) carriers who may not experience clinical symptoms for many years. The high rate of progression of acute infection to chronic infection (70-100%) and liver disease (>50%), its world-wide distribution and lack of a vaccine make HCV a significant cause of morbidity and mortality.

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HCV is an enveloped virus whose genome is a 9.4kb single-stranded RNA (sense(+)) encoding a single polyprotein that is processed by proteolysis to yield at least 9 proteins. HCV is related to pestiviruses and flaviviruses (Choo, Q-L. et al. 5 (1989) *Science* **244**:362-364; Choo, Q-L. et al. (1991) *Proc. Natl. Acad. Sci. USA* **88**:2451-2455. Reinfection of previously HCV-infected chimpanzees suggests that protective immunity is transient or non-existent (Farci, P. et al (1992) *Science* **258**:135-140). Furthermore, results of recent vaccine trials 10 suggest that development of an effective vaccine is remote (Houghton, M. et al. (1994) *2nd Internat. Meeting on Hepatitis C (San Diego)*). Attempted treatment of chronic HCV infection using existing antiviral agents produces low cure rates and serious side effects. (Dienstag, J.L. (1983) *supra.*)

15 The nucleotide sequence of the HCV genome has been cloned and a single open reading frame has been identified. Using a vaccinia virus expression system, several cleavage products have been tentatively identified. (Lin, C. et al. (1994) *J. Virol.* **68**:5063-5073; Grakoui, A. et al. (1993) *J. Virol.* **67**:1385-1395.) The various putative cleavage products were recognized by antibodies raised against various peptides synthesized from amino acid sequences deduced from various segments of the coding regions. Sizes of antibody-reactive peptides were estimated by 20 SDS-PAGE (See Fig. 1). The non-structural protein designated 5B (NS5B) has been shown to have an amino-terminal sequence SMSY (Ser-Met-Ser-Tyr). The NS5B region encodes a 68kd protein (p68) which contains an internal GDD (Gly-Asp-Asp) motif found in RNA-dependent RNA polymerases of other RNA viruses (Koonin, E.V. 25 (1991) *J. Gen. Virol.* **72**:2197-2206). However, no polymerase activity has been detected for HCV p68. In fact, the question has been raised that the 5B protein (p68) alone does not encode an active RNA-dependent RNA polymerase enzyme and that another 30 subunit, possibly the NS5A gene product, is essential to catalytic activity. Prior attempts by the inventors and others to express the NS5B coding region as a fusion protein, using 35 existing expression systems that facilitate purification of the

fusion product, using existing expression systems that facilitate purification of the fusion product and specific cleavage have failed to yield any active polymerase.

5 Summary of the Invention

The present invention provides a recombinant protein of HCV having RDRP activity (r-HCV-RDRP) obtainable by expression in a host mammalian or bacterial cell of a modified NS5B coding region of HCV. The modification includes addition at the amino terminus 10 of a methionine residue and optionally from 1-20 additional amino acids interposed between the N-terminal methionine and the N-terminal serine of unmodified NS5B gene product. The modification also includes deletion at the amino terminus of up to 9 amino acids to provide an amino-terminal methionine. Two 15 methionines occur naturally according to the deduced sequence of wild-type HCV-RDRP. Therefore, modification includes deletion to remove amino acids lying N-terminal to either methionine or, alternatively, deletion to some intermediate point between the two methionines plus addition of an N-terminal methionine codon. 20 A combination of deletions and insertions, within the limits described is also contemplated. Added amino acid sequence can be devised to create a specific protease cleavage site to permit post translational modification of the recombinant HCV-RDRP expression product, *in vivo* or *in vitro*. Such post- 25 transcriptional modification can be used to generate exactly the amino acid sequence encoded by NS5B, having an N-terminal serine. Added amino acid sequence can be devised to generate an affinity ligand binding site, for convenience and ease of purification. The data reported herein were obtained with a r-HCV-RDRP having 30 an N-terminal MA (Met-Ala) dipeptide, giving an N-terminal sequence MASMSY (SEQ ID NO:6) instead of the predicted SMSY sequence of the wild-type processed protein. The coding sequence of NS5B is accordingly modified to include a met codon (ATG) at the 5'-end, as well as, optionally, codons for other amino acids 35 to be included or deleted. Minimal modifications are preferred, in order to avoid potential deleterious effects on enzyme activity, and to avoid creating artificial epitopes. The r-HCV-

RDRP can be expressed in *E. coli* and in mammalian cells to yield active RDRP. The expression of active r-HCV-RDRP in *E. coli* demonstrates that no other HCV-encoded protein is necessary for polymerase activity.

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The invention further provides r-HCV-RDRP in solubilized form, and a method of solubilization without destroying activity.

10 The invention also provides methods for purifying solubilized HCV-RDRP. One such method, to be used in combination with others, is affinity chromatography, using antibody to r-HCV-RDRP as the affinity ligand. Other affinity ligands are obtained by a combinatorial library approach as described, e.g., by Wu, J. et al. (1994) *Biochemistry* 33:14825-14833; and Ohlmeyer, 15 M.H.J. et al. (1993) *Proc. Nat. Acad. Sci. USA* 90:10922-10926.

20 In addition, the invention provides polyclonal or monoclonal antibodies specific for HCV-RDRP. Such antibodies can be made by known techniques, using the purified enzyme as antigen. Such antibodies bind either r-HCV-RDRP or wild-type HCV-RDRP. The availability of such antibodies makes it possible to prepare an affinity-labeled chromatography matrix for rapid purification of HCV-RDRP. The antibody also makes possible rapid detection of HCV-RDRP in biological materials, for example, in serum of HCV-infected patients.

25 The invention further provides a method for transfecting a mammalian cell with HCV-RDRP and expressing the enzyme within the cell. Consequently, the invention also provides a transfected mammalian cell line expressing r-HCV-RDRP. Such cells are useful for assaying the effects of candidate anti-viral compounds as inhibitors of RDRP activity.

35 Therefore, the invention also provides a method for screening possible inhibitors of RDRP activity *in vivo*. Compounds with inhibitory activity can have anti-viral activity, since inhibition of the polymerase inhibits viral replication and

expression of virus gene products. The *in vitro* assay is advantageous because it can rule out compounds which cannot enter the infected cell. One class of attractive candidate compounds is the nucleoside analogs; compounds which after being modified (phosphorylated) within cells can bind to substrate sites on the enzyme or which can be incorporated into a newly synthesized RNA but whose presence there disrupts normal function of the HCV polymerase or further replication of an RNA containing the analog. Acyclovir is one example of a very effective and safe nucleoside analogue that inhibits DNA virus replication by inhibiting a viral polymerase (DNA-dependent DNA polymerase) and interfering with primer-template function (chain termination). Such analogs are almost always effective only in the nucleotide triphosphate form. The *in vitro* assay provides a convenient method of administering the compound in its nucleoside form or nucleoside monophosphate form, allowing endogenous metabolic activity of the cell to convert that form to the active triphosphate, thereby avoiding a step of chemical synthesis of the triphosphate, as would be required for an *in vivo* assay.

20 A method for measuring HCV-RDRP activity *in vitro* is also provided. Such an assay permits identification of the enzyme and evaluation of its concentration during purification. In addition, the assay provides an additional, *in vitro*, method for screening potential inhibitors of RDRP as candidate anti-viral agents.

30 In principle, any compound can be tested as a candidate RDRP inhibitor. Certain classes of compounds are considered attractive candidates. These include, without limitation, nucleoside analogs, oligonucleotides and peptides. Certain compounds having planar, polycyclic-aromatic characteristics are also potential inhibitors. It will be understood that compounds identified as effective RDRP inhibitors must be further screened for toxicity, bioavailability, side effects and the like before being tested as therapeutic agents. Nevertheless, the initial identification as an inhibitor of HCV-RDRP is an essential first

step in the development of an anti-viral therapy. It will also be recognized that an inhibitor of r-HCV-RDRP will also inhibit wild-type HCV-RDRP.

5        In another aspect of the invention, the existence of purified HCV-RDRP or r-HCV-RDRP makes it possible to detect and measure antibodies to RDRP present in the serum of an HCV-infected patient. The fact that such antibodies exist at all is in itself a finding made possible by the expression and preparation of purified r-HCV-RDRP according to the invention. The existence of circulating antibodies to HCV-RDRP in infected serum may be due to lysis of infected cells and release of HCV-RDRP into the extracellular fluids and bloodstream, where it can stimulate an antibody response. As the disease fluctuates in 10 severity, the amounts of HCV-RDRP released and the amounts of antibody thereto would also fluctuate. Therefore, the amount of antibody to HCV-RDRP present in a patient's serum can be used as an indicator, not only of the presence of infection, but of its 15 severity at a given time. The assay for anti-HCV-RDRP can serve as a means of diagnosing infection and also as a means of monitoring the course of the disease over time or in response to treatment. The assay for anti-HCV-RDRP can be carried out by a 20 variety of known techniques, such as the gel separation method described herein. Other suitable methods include ELISA, and 25 radioimmunoassay. A sandwich-type assay, using immobilized r-HCV-RDRP to capture the antibody can then use an anti-immunoglobulin reagent tagged with an appropriate marker such as an enzyme, radioisotope, fluorescent molecule or chemiluminescent marker or the like, all as understood by those skilled in the 30 art. (Antibodies: A laboratory manual, Ed Harlow and David Lane, Cold Spring Harbor Laboratory (1988) pp. 553-611.)

#### Brief Description of the Figures

35        Fig. 1. Hepatitis C virus genome and polyprotein cleavage products. The cleavage products of the HCV polyprotein have been tentatively identified using vaccinia virus expression systems. The amino terminus of the 5B protein expressed and processed in

this system is SMSY (Ser-Met-Ser-Tyr). Although published reports have not proved that the 5B protein has RNA polymerase activity, it does contain the GDD (Gly-Asp-Asp) motif found in other RNA-dependent RNA polymerases. The question has been raised that the 5B protein alone does not encode an active RNA-dependent RNA polymerase enzyme and that another subunit (possibly the NS5A gene product) is essential for catalytic activity. During the initial phases of this work we were unsure if the protein encoded by NS5B would exhibit RNA-dependent RNA polymerase activity simply due to the lack of other essential factors.

Fig. 2. Expression of r-HCV RNA-dependent RNA polymerase in *E. coli* using the T7 polymerase driven Studier vectors. *E. coli* containing the engineering T7 polymerase driven expression vector were incubated at 37°C until an OD<sub>600</sub> of 0.6 was reached. A sample of cells was obtained and IPTG added to a final concentration of 1 mM. Samples were collected at 1, 2 and 3 hours after IPTG induction. Whole cells were lysed in 1X sample buffer at 95°C and samples analyzed by 10% SDS-PAGE. The photograph shows a representative Coomassie Blue stained gel. Lane 1 represents molecular mass markers; lane 2, the uninduced control (0 h); lane 3, 1 h; lane 4, 2 h; and lane 5, 3 h after IPTG induction. Recombinant r-HCV RNA-dependent RNA polymerase is indicated by an arrow (RDRP).

Fig. 3. Some patients with chronic hepatitis C have circulating antibodies that react with recombinant HCV RNA-dependent RNA polymerase. Cells expressing r-HCV RDRP were harvested and lysed by heating in SDS-PAGE sample buffer. Soluble proteins were separated by SDS-PAGE, transferred to nitrocellulose membranes and immunoblotted with human sera using an Immunetics Miniblitter template (Hagedorn, et al. *FEBS Lett.* (1990) 264:59-62). Immunoblots were developed with a secondary anti-human horseradish peroxidase conjugated antibody and enhanced chemiluminescent methods (ECL, Ambersham). This photograph shows an immunoblot where lane 1 was probed with

normal human serum and lanes 2 (1:500 dilution), 3 (1:300), and 4 (1:200) were probed with serum from a patient with chronic hepatitis C. The location of recombinant HCV RDRP (visualized by Coomassie and Ponceau S staining) is indicated by an arrow. 5 These lower molecular mass bands seen in lanes 3 & 4 represent proteolytic fragments of RDRP seen when whole *E. coli* lysates are used in immunoblots.

Fig. 4. Solubilization of HCV RDRP under nondenaturating conditions. Cells expressing r-HCV RDRP were harvested and processed using standard methods. Samples of insoluble *E. coli* pellets (pellet, positive control) and soluble fractions from cells containing or not containing (negative control) the RDRP expression vector were separated by SDS-PAGE and transferred to nitrocellulose membranes. Proteins bound to nitrocellulose were probed with rabbit preimmune and rabbit anti-RDRP sera as outlined in Fig. 3. Blots were developed with the ECL system (Amersham). The photograph shows an immunoblot where lanes 1, 3, 5, 7 and 9 were probed with preimmune serum and lanes 2, 4, 6, 8 and 10 were probed with immune serum. Two independently prepared samples of soluble proteins (plus IPTC #1 & #2, lanes 3-6) were examined.

Fig. 5. Enzymatically active recombinant r-HCV RNA-dependent RNA polymerase. Poly(U) polymerase activity of purified recombinant poliovirus RDRP (approximately 50 ng) and equal quantities of soluble protein (approximately 1  $\mu$ g) from *E. coli* expressing r-HCV RDRP (HCV RDRP lysate) or control cells not expressing RDRP (control lysate) are shown. Incubations were performed as described previously and CPM of poly(U) recovered from 15  $\mu$ l of incubation are shown at 30 min (one sample) and 60 min (mean of duplicates) of incubation are shown. (J. Virol. 30 (1986) 58:790-796).

Fig. 6. Enzyme assay of partially purified recombinant HCV RNA-dependent RNA polymerase. Lysates of *E. coli* expressing r-HCV RDRP were prepared and enzyme assays performed as in Fig. 5.

The experiment shown used a DEAE resin in a batch purification approach under pH and buffer conditions that allow RDRP to bind the resin. Equal quantities of protein were assayed from cell lysates (designated RDRP lysate) and proteins eluted from DEAE resin with 0.5 M NaCl (designated DEAE batch) that were concentrated to approximately that of the lysate. Additional studies with DEAE resin and other test resins have shown that partial purification of RDRP enzyme activity correlates with increases in the amount of unproteolyzed RDRP that we detect by immunoblotting using rabbit antiserum and the methods outlined in Fig. 4.

Fig. 7. Immunoaffinity purification of recombinant r-HCV RDRP. This photocopy shows an initial result with an immunoaffinity column prepared with rabbit polyclonal protein A Sepharose purified anti-RDRP antibodies. The starting material was protein solubilized from *E. coli* expressing recombinant r-HCV RDRP. The figure shows an immunoblot of proteins that were eluted from two identical columns that had protein applied under different detergent conditions. Lane 1 depicts proteins eluted from antibody/Sepharose beads that had been mixed overnight with the solubilized recombinant RDRP in 20 mM Tris-pH 7.5, 100 mM KCl, 0.5 mM EdTA, 1 mM DTT, 5% glycerol, and 0.05% Triton X-100. These beads were washed the following morning with 10 mM potassium phosphate buffer-pH 7.2 and proteins eluted with 100 mM glycine-pH 2.5. The eluted protein was collected in 1 M Tris-pH 8.0 to readjust the pH. Proteins eluted from the column were then analyzed by SDS-PAGE and immunoblotting as in Fig. 4. Lane 2 depicts proteins eluted from identical beads mixed with the same starting material except that 0.05% NP-40 was present instead of 0.05% Triton X-100. The location of r-HCV RDRP is indicated.

Detailed Description of the InventionDefinitions

"RDRP" stands for RNA-dependent RNA polymerase, an enzyme catalyzing RNA synthesis, the synthesized RNA having a sequence complementary to an RNA template. "HCV-RDRP" is the RDRP of Hepatitis C virus. The modified HCV-RDRP described herein is designated r-HCV-RDRP. The region of the HCV genome designated NS5B has been identified as a protein cleavage product of the HCV polyprotein, using a vaccinia virus expression system as described *supra*. The nucleotide sequence of NS5B is included in SEQ ID NO:1. Putative amino acid coding by the NS5B sequence begins with nucleotide 7. Where the sequence has been deleted at the 5' end, the remaining sequence has been designated by the nucleotide numbers beginning and ending the remaining coding sequences, not including the stop codon. For example NS5B<sub>34-1779</sub> designates that part of NS5B including nucleotides 34-1779 of SEQ ID NO:1.

The amino acid sequence encoding the r-HCV-RDRP exemplified herein is given in SEQ ID NO:2. The amino acid sequence encoded by NS5B begins at amino acid No. 3 of SEQ ID NO:2. Where the sequence encoded by NS5B has been deleted at the N-terminus, the remaining sequence is designated  $\Delta^n$ NS5B where n is the number of amino acids deleted from the N-terminus of NS5B. For example  $\Delta^9$  NS5B is the sequence of amino acids 12-593 in SEQ ID NO:2.

Various r-HCV-RDRP constructs are contemplated according to the invention, as described. Modified coding sequences included with the invention have the general sequence

ATG-N<sub>x</sub>- (N<sup>S</sup>) (N<sup>M</sup>) (N<sup>S</sup>) (N<sup>Y</sup>) (N<sup>S</sup>) (N<sup>W</sup>) (N<sup>T</sup>) (N<sup>G</sup>) (N<sup>A</sup>) - [NS5B<sub>34-1779</sub>] where N<sub>x</sub> is any nucleotide sequence encoding from 0-20 amino acids, N<sup>S</sup> is a codon encoding serine, N<sup>M</sup> is a codon encoding methionine, N<sup>Y</sup> is a codon encoding tyrosine, N<sup>W</sup> is a codon encoding tryptophane, N<sup>T</sup> is a codon encoding threonine, N<sup>G</sup> is a codon encoding glutamic acid, and N<sup>A</sup> is a codon encoding alanine. Any of the codons in parentheses can be deleted, if desired. Up to 5 of the codons in parentheses can be mutated if desired. The term "mutated" is

intended to mean altered to encode an amino acid other than that originally encoded by the NS5B sequence. For example, individual codons can be altered to encode alanine, by the known method of alanine scanning mutagenesis. Alanine scanning mutagenesis provides a rapid and convenient method for identifying amino acid positions where substitution is tolerated, without substantially affecting function negatively. Positions where alanine scanning reveals tolerance for substitution are likely to tolerate other amino acid substituents as well. Preferred substituents are one or more histidine residues, which can serve as affinity ligands for metal (e.g. nickel) columns. The presence of histidine provides preferential binding to the column to facilitate purification of r-HCV-RDRP. [NS5B<sub>34-1779</sub>], as defined, represents the nucleotide sequence of the remainder of SEQ ID NO:1, from nucleotides 34-1779, not including the stop codon. The techniques for making any of the foregoing sequences are essentially as described below for the sequence where N<sub>x</sub> is GCT and none of the codons in parentheses, encoding the first nine amino acids encoded by NS5B, is deleted. It will be apparent that primers can be synthesized for the desired sequence combined with desired restriction site sequences to facilitate insertion into appropriate expression vectors. The choice of vector is based on factors known in the art, including the host cell, the type of promoter desired and the presence or absence of additional sequences which could be co-expressed with the r-HCV-RDRP. The reaction condition, PCR, vector insertion and host cell growth are as described below or as well-known in the art.

Further modifications can be made to r-HCV-RDRP. A deletion of approximately 25% of the C-terminal region that did not include the Gly-Asp-Asp motif was constructed (r-HCV-RDRP-ΔC) and tested for activity. Although protein was expressed and was detectable by immunoblotting with anti-RDRP serum, lysate of *E. coli* expressing r-HCV-RDRP-ΔC had no measurable activity in either the poly(U) assay or with globin mRNA as template. However, the region of amino acids 565-572 of NS5B (565-574 of SEQ ID NO:2) near the C-terminus is considered to be exposed at

the protein surface on the basis of modeling studies. Site directed mutagenesis has been used to alter the coding to Arg 570-His, Arg 572-His and Try 573-His (using the numbering of SEQ ID NO:2). By clustering one or more additional histidine residues in a surface region near one end of the protein, purification by metal-affinity chromatography is facilitated while enzymatic activity is not substantially affected. Other predicted surface regions that can serve as sites for mutagenesis to replace an existing amino acid with histidine include amino acids 47-56, 152-159, 183-184, 210-215, 269-272, 384-391, and 43999-442 in SEQ ID NO:2.

Although the r-HCV-RDRP can be expressed in virtually any host cell type, the enzyme should preferably be in soluble form in order to be useful in *in vitro* studies, such as testing for inhibitors. When synthesized intracellularly, the enzyme is in an insoluble form in cell lysates unless steps are taken to solubilize the enzyme. In general, host cells are collected and concentrated, then lysed by means known to disrupt the host cells, for example by the use of a host-cell-wall hydrolyzing enzyme, by sonication and the like. In general a protease inhibitor is added to protect against proteolytic enzymes released by cell lysis. A non-ionic detergent can also be employed. Sonicated cells and subcellular complexes are subjected to freezing and thawing in the presence of the above-named components. Remaining particulate matter is removed by centrifugation at 10,000-35,000xg. The r-HCV-RDRP remains in the supernatant. A detailed protocol for solubilizing r-HCV-RDRP expressed in *E. coli* is described below.

Further purification of the enzyme is accomplished by techniques and expedients known in the art. These include, but are not limited to, antibody affinity chromatography, metal-binding affinity chromatography (a technique especially suited for modified forms of the enzyme having added histidine residues) as well as conventional ion-exchange columns, differential precipitation with ammonium sulfate and other methods known in

the art, not limited to the methods specifically disclosed herein. "Purified form" is used herein to mean any preparation of the enzyme having at least 4-fold greater specific activity than that measured in a solubilized cell lysate.

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Antibodies against r-HCV-RDRP can be generated by a variety of known techniques for making monoclonal or polyclonal antibodies. Antibodies to r-HCV-RDRP also bind to HCV-RDRP of infected cells, as shown by the discovery that circulating antibodies to HCV-RDRP are detectable in serum of HCV-infected patients, using r-HCV-RDRP as the antibody ligand. A variety of monoclonal antibodies can be selected, having affinity for different epitopes of r-HCV-RDRP, as known in the art. Some antibodies can be inhibitory of enzyme activity. Others can have a modest affinity that facilitates binding to an antibody-affinity column and subsequent elution under conditions that do not inactivate the enzyme.

Mammalian cells are a preferred host cell for certain purposes, particularly for *in vitro* screening for inhibitors of HCV-RDRP, and also for developing cell lines that can propagate HCV in cell culture. Any of the known cell lines used for transformation can, in principle, be transformed to express r-HCV-RDRP. Preferred cell lines are those of tissue origin known to be infected by HCV or similar viruses, such as flaviviruses. Such cell lines include, for example, the human macrophage cell line U937, human liver-derived hepG2 cells, and the pig kidney cell line PK15. A recently-discovered segment near the 3'-end of HCV-RNA has a predicted cloverleaf-type secondary structure that is a likely site of interaction for a protein or peptide (which may be host-encoded) that regulates HCV-RDRP (Tanaka, T., et al. (1995) *Biochem. Biophys. Res. Commun.* 215:744,749). Such regulation can take the form of altering the template specificity or the catalytic activity of r-HCV-RDRP as well as the wild-type enzyme. Expression of r-HCV-RDRP in cells that normally express the regulatory protein provides an *in vivo* cell system where expression of r-HCV-RDRP closely approaches the manner of

expression in HCV-infected cells. Also, the presence of active HCV-RDRP in a cell can enhance the replication of HCV introduced by infection or genomic HCV RNA introduced by transfection, by providing a "jump start" for HCV replication. Most importantly, 5 the ability to measure r-HCV-RDRP activity in transformed cells provides an essential key to screening potential inhibitors of HCV-RDRP for their ability to inhibit the enzyme *in vivo*.

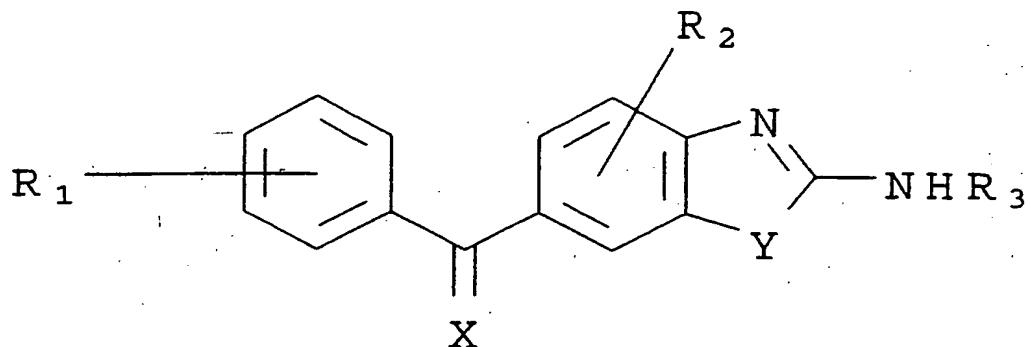
10 Assay for RNA synthesis by r-HCV-RDRP *in vitro* has demonstrated that the enzyme can catalyze synthesis of poly(U) using a poly(A) template, and synthesis of RNA, using a globin mRNA template. Both reactions were >90% primer-dependent under the described reaction conditions. The enzyme had no detectable activity in the absence of Mg<sup>++</sup> and showed maximal activity in 15 the presence of about 20 mM Mg<sup>++</sup>. A method for screening compounds that have anti-viral activity is provided by testing the effect of various compounds on the RDRP enzyme activity *in vitro*. The *in vitro* method includes comparing the amounts of RNA synthesized in the presence and absence of a test compound. 20 An inhibitory effect is indicated if the amount of RNA synthesized is reduced in a reaction where the test compound is present compared to a control reaction where the test compound is absent.

25 The stably transfected cell line expressing r-HCV-RDRP is especially useful for carrying out *in vivo* screening for compounds that inhibit the polymerase in intact mammalian cells. Such inhibitors are likely inhibitors of HCV replication in infected cells. An *in vivo* test is advantageous since it can 30 screen out cytotoxic compounds, and compounds that cannot enter the cell in sufficient quantity. It also provides the added capability of testing compounds which are precursors of the actual inhibitor but converted to inhibitor by the cells metabolic processes. Examples of precursor inhibitors include 35 nucleoside analogs which must be phosphorylated to be converted to active inhibitor, and compounds having an active group protected by a linkage that undergoes hydrolytic cleavage,

intracellularly, to form active inhibitor. Non-nucleoside analogs having the general structure shown are considered attractive candidates for r-HCV-RDRP inhibitors

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where R<sub>1</sub> and R<sub>2</sub> are alkyl, amino, hydroxy, alkoxy or halo groups, R<sub>3</sub> is an alkyl, aryl, hydroxy or alkoxy group, X is O, NR<sub>3</sub>, O, CH<sub>2</sub> or CHR<sub>3</sub>. In general, the test cells additionally are transformed with a reporter construct whose expression requires the action of r-HCV-RDRP, or whose expression is amplified by the presence of r-HCV-RDRP. Reporter genes are well known in the art, including, but not limited to luciferase, secreted alkaline phosphatase and the fluorescent green protein, all of which are commercially available. An attractive strategy is to use an antisense gene for the reporter, that is, a version of the reporter gene which expresses an antisense, or (-) strand messenger RNA of the reporter gene. Activity of an RDRP is then required to produce a sense (+) strand in RNA which can be translated to yield active reporter. This system has the advantage that there is no background level of reporter activity in the absence of active HCV-RDRP, if the RDRP is inhibited. An outline for the construction of suitable (-) strand reporter gene is set forth below.

#### Material and Methods

Material - All chemicals were purchased from Fisher and all enzymes from Gibco BRL unless stated otherwise. AmpliTaq was

5 purchased from Perkin-Elmer. All other PCR and ligation components were from Invitrogen. Lysozyme, antibiotics, and pre-stained protein standards were from Sigma. Nucleotides and poly(A) were from Pharmacia. [<sup>3</sup>H-]UTP was from Dupont NEN. Oligo(U) was a generous gift from E. Ehrenfeld (University of California, Irvine).

10 Subcloning of the HCV NS5B region - PCR-primers for the amplification of the NS5B-region were designed based on the N-terminus as predicted by vaccinia virus expression studies (Lin, C. et al. (1994) *supra*; Grakoui, A. et al. (1993) *supra*) and the C-terminus based on the end of the open-reading-frame of the HCV poly protein (Choo, Q-L. et al. (1991) *supra*). The template was the original prototype HCV (type 1a) clone (obtained from the 15 CDC) (Choo, Q-L. et al. (1989), (1991), *supra*). Using the following primers, 5'-ATA GCT AGC ATG TCT TAC TCT TGG ACA GG-3' (SEQ ID NO:3) and 5'-ATA GGA TCC TCA TCG GTT GGG GAG GAG G-3' (SEQ ID NO:4), we amplified the NS5B-region with minimum changes at the N-terminus (ASMSY SEQ ID NO:5 instead of SMSY SEQ ID NO:7) and 20 directionally cloned it into pET-11a (Novagen) at NheI and BamHI restriction sites (Maniatis et al. (1982) Molecular Cloning, Cold Spring Harbor Laboratory, Plainview, New York). The PCR amplified gene had a NheI site engineered into the 5' end and a BamHI site at the 3' end. This construct results in the 25 synthesis of a recombinant protein with an amino terminal sequence of MASMSY rather than the SMSY amino terminus of the putative wild-type NS5B protein predicted by vaccinia virus expression studies.

A S M S Y S W T (amino acid encoded in primer, SEQ ID NO:8)

5'-ATA GCT AGC ATG TCT TAC TCT TGG ACA GG-PCR primer directed at the  
 ↑  
 5 NheI cut site  
 10 C-A  
 A-C  
 C-G  
 C-G  
 C-G  
 A-T  
 G-T  
 A-T  
 G-C  
 G-C +26  
 15 pppG-C CTAGAAATATTTTCTT  
 20 PET-11a +50 +61 Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser  
 25 SD NheI HheI +50 +61 Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser  
 30 BamHI ISEQ ID NO:10 ISEQ ID NO:91

This region of the expression vector is removed before inserting PCR product

25 Restriction site in the pET-11a expression vector where the 5' end of the PCR amplified NS5B gene was inserted.

NheI  
 G>CTAGC

30 The PCR reaction was preceded by an 2 min incubation at 94°C, followed by 20 cycles of 1 min 94°C, 2 min 55°C, and 3 min 72°C (Coy Corporation Tempcycler II). Reactions were completed by keeping the temperature at 72°C for another 7 min and subsequent cooling to 4°C. Amplified DNA was purified by phenol/chloroform extraction, digested with NheI and BamHI and 35 re-purified using phenol/chloroform extraction and ethanol precipitation.

40 Insert and vector were ligated overnight at 14.5°C at an 3:1 ratio (insert:vector) using T4-ligase (Maniatis, *supra*). Ligated material was used to transform *E. coli* (Top10™ from Invitrogen) using CaCl<sub>2</sub> methods. Colonies were selected on ampicillin plates and minipreps of plasmid DNA isolated from single colonies were

characterized using restriction enzyme analyses. Plasmid DNA obtained by mini-preparation methods was used to transform BL21 (λDE3) *E. coli* (Novagen), organisms containing plasmid were selected using ampicillin and mini-preparations of plasmid DNA from single colonies were analyzed by restriction enzyme digestion.

10 *Expression, purification and solubilization of the putative HCV RDRP - BL21 (λDE3) E. coli* containing the pET-11a-NS5B construct described above were grown in overnight cultures (M9ZB media with Carbenicillin) and diluted 1:20 into fresh medium the next morning. Cells were incubated at 37°C until the culture media reached an OD<sub>600</sub> of 0.6. IPTG was added at that time to a final concentration of 1 mM. Expression of the putative RDRP was 15 followed by SDS-PAGE analysis of whole cells lysed in sample buffer at 90°C.

20 To solubilize RDRP under non-denaturing conditions cells were harvested 2h after IPTG-induction. RDRP was solubilized by lysing the cells on ice for 20 min in 20 mM Tris pH 7.5, 100 mM KCl, 0.5 mM EDTA, 1 mM DTT, 0.1% Triton X-100 and 30 µg/ml lysozyme. Samples were sonicated on ice with an 0.5 inch probe (pulse setting) for 5 min (Ultrasonics Inc. W-225, output-setting 7) and 25 centrifuged (19,000 g at 4°C for 30 min). The insoluble fraction (pellet) obtained from these preparations was enriched with RDRP. Pellets were suspended in SDS-PAGE sample buffer and heated for 10 min at 90°C and used as RDRP markers for SDS-PAGE gels. However, active enzyme was found in the supernatant, as set forth in the following protocol.

30

Outline of Solubilization Method for Recombinant RDRP

1. Thaw 5 g of *E. coli* pellet.

35

2. Resuspend 5 g of pellet in:

- 45 ml Lysis buffer
- 40  $\mu$ l 100mM PMSF (plus other protease inhibitors)
- 150  $\mu$ l lysozyme (10 mg/ml)

5

Lysis buffer:

20 mM Tris pH 7.5 (at 4°C)  
0.5 mM EDTA  
100 mM KCl  
1 mM DTT  
0.1% Triton X-100 (or 0.1% NP-40)  
10.0% (v/v) Glycerol

10

15

3. Place samples on ice for 20 min, then sonicate for 5 min (pulse mode; setting between 6-7). Mix while sonicating.

4. After sonicating flash freeze the lysate in liquid nitrogen (put the lysate into liquid nitrogen for about 1-2 min).

20

5. Quickly thaw the lysate at 37°C water bath.

6. Sonicate the lysate for 1 min.

25

7. Add an additional 5 ml of lysis buffer per 45 ml of sonicated sample mix.

30

8. Divide entire sonicated sample into 50 ml fractions (Fisher 50 ml tubes).

35

9. Centrifuge lysate at 12,500 rpm for 20 min in Beckman J-17 rotor (or 12,500 rpm in a Sorvall SS-34 rotor).

40

10. Remove supernatants to clean (sterile) 50 ml Fisher tubes and add sterile protein grade glycerol to a final concentration of 10% (for example, 4.44 ml of glycerol/40 ml of supernatant). This solution is stored at 4°C and used as starting material for the purification of enzymatically active HCV RDRP.

45

Further purification is accomplished by employing the following steps, either singly or in combination.

**Soluble proteins from lysate of *E. coli* expressing recombinant HCV RDRP  
(10,000 x g supernatant)**

↓

**5 45% ammonium sulfate precipitated proteins  
(subsequently dialyzed in Tris-pH 7.5  
with 10% glycerol and 1mM DTT)**

↓

**10 DEAE anion exchange chromatography  
(starting material applied at pH 8.5 - Tris buffer,  
wash step with same buffer and enzyme  
is eluted with a 0-0.5 M NaCl gradient)**

↓

**15 Phosphocellulose chromatography  
(concentrated DEAE fractions containing enzyme  
are diluted to decrease salt concentration and change  
pH to 8.0 (Tris/HCl) before applying samples to column,  
wash step, and elution with a 0-0.5 M NaCl gradient)**

↓

**20 25 FPLC gel filtration chromatography  
(Superose 12 HR 10/30 - Pharmacia)  
(starting material is concentrated, buffer changed  
to 150 mM NaCl with Tris pH 8.0 and applied in  
a 200  $\mu$ l volume with a flow rate of 0.3 ml/min)**

↓

**30 Recombinant HCV RDRP for enzyme assays  
(stored in aliquots at -70°C with Tris-pH 8.0,  
100 mM NaCl, 20% glycerol, 0.1% NP-40, & 1 mM DTT)**

**35 40 As in all protein purification procedures, one can modify  
buffers, pH and other conditions to further optimize the  
purification of HCV RDRP. An additional final purification step  
(or substitution for the FPLC gel filtration step) is a Mono-S  
cation exchange chromatography step at pH 6.0 with a MES buffer  
(the isoelectric point of the enzyme is approximately 8.8). All  
purification steps are monitored for enzyme activity using RDRP  
assay, total protein, and analyzed by SDS-PAGE.**

**45 Rabbit anti-HCV RDRP serum - RDRP solubilized from the pellet  
fraction as described above was separated by preparative SDS-PAGE  
and used to immunize rabbits. Animals were immunized at 4-5 week  
intervals as described in detail previously. (Harlow, E. and D.**

Lane (1988) Antibodies: A laboratory manual, Cold Spring Harbor Laboratory, pp. 553-611.)

5        *Immunoblotting analysis* - Immunoblots were performed using previously described methods with the modification that secondary HRP conjugated antibodies were used with the enhanced chemiluminescent system (ECL, Amersham). When rabbit serum was the primary antibody, the secondary antibody was anti-rabbit immunoglobulin. When human serum was screened the secondary antibody was anti-human immunoglobulin. Serum from patients with documented chronic hepatitis C infections was provided by Dr. Michael Beach of the Centers for Disease Control and Prevention (Atlanta).

15       *Poly(U) polymerase assay* - Enzyme activity in soluble fractions was measured using a poly(U) polymerase assay with poly(A) as template and oligo(U) as a primer (Hey, T.D. et al. (1986) *J. Virol.* **58**:790-796). Samples (generally 2  $\mu$ l) were assayed in 50  $\mu$ l incubations containing 50 mM HEPES (pH 8.0), 500  $\mu$ M each ATP, CTP, and GTP; 4 mM DTT, 3 mM MgAc<sub>2</sub>; and 60  $\mu$ M ZnCl<sub>2</sub>. [<sup>3</sup>H]UTP at a concentration of 15  $\mu$ M (specific activity: 27 Ci/mol) was also present. Each incubation contained 1  $\mu$ g of poly(A) and 0.5  $\mu$ g oligo(U) as a primer.

25       Incubation were at 30°C for 30-60 min, [<sup>3</sup>H]poly(U) was precipitated with TCA in the presence of carrier DNA and collected on Whatman GF/C filters. Filters were washed with 0.1 M sodium pyrophosphate/1 N Hydrochloric acid and 95% ethanol, respectively. [<sup>3</sup>H]poly(U) was quantitated by liquid scintillation spectrometry (LKB 1218 RackBeta).

Stable Transfection of Baby Hamster Kidney (BHK)  
Cells Using Lipofectin

Day 1 (Afternoon)

5

Split the BHK cells into 6 well plates aiming for 50% confluence for transfection

Day 2 (after 4 p.m.)

10

Prepare the following solutions in sterile tubes:

(A) 50  $\mu$ l miniprep DNA + 50  $\mu$ l media without serum (DMEM/F12)  
(2 each)

15

(B) 6.25  $\mu$ l Lipofectin (Life Technologies, Gaithersburg, MD) +  
93.75  $\mu$ l media

20

(C) 12.5  $\mu$ l Lipofectin + 87.5  $\mu$ l media

(D) 6.25  $\mu$ l Lipofectin + 193.75  $\mu$ l media (mock transfection)

(E) 12.5  $\mu$ l Lipofectin + 187.5  $\mu$ l media (mock transfection)

25

Gently mix A&B and A&C and let the DNA and Lipofectin react for 15 minutes at room temperature. During this time, wash the cell twice with 2 mls of DMEM/F12. Add 1.8 mls of DMEM/F12 to the DNA/Lipofectin complex and add it to the cells with gentle swirling. Leave the cells in the incubator overnight.

30

Day 3 (9 a.m.)

35

Remove the DNA/Lipofectin and add 3 mls of media + serum to the cells. Incubate the cells for 30-48 hours. Split the cell 1:20, 1:50 and 1:100 into 10 cm dishes in 10 mls of media + serum containing 600  $\mu$ g/ml geneticin. Allow 3-7 days for selection and 10-14 days for colony formation. The same protocol can be adapted to employ Starburst Dendrimer (Life Technologies,

Gaithersburg, MD) instead of Lipfectin, to improve transfection efficiency.

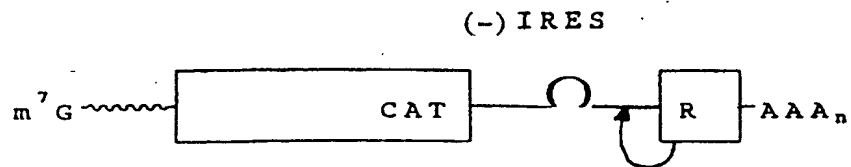
5 After selection, ring clone colonies onto 24 well plates and assay media from confluent wells for RDRP activity. Maintain cells n 600  $\mu$ g/ml geneticin.

10 Use of Stably Transfected Cells Expression HCV RDRP to Identify Compounds that Enter Intact Cells and Inhibit HCV RDRP

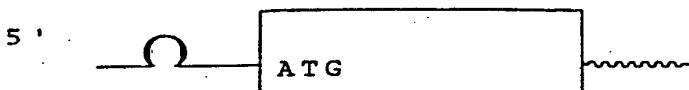
15 The most direct approach to determining the effect of potential inhibitors of HCV RDRP in transformed cells is to directly measure RDRP activity in cell extracts after cells have been incubated with compounds and washed extensively. This can be done using the RDRP assay described herein (with a HCV template) and requires no other new development except the cell-line. In brief, cells are incubated under conditions that maximally express active enzyme and in sufficient quantities for subsequent for subsequent enzyme assays. Test compounds are 20 added to incubations, media are removed at the desired time and cells are extensively washed to remove extracellular test compounds. Extracts of cells are prepared for RDRP assays following the general methods described herein. This approach is relatively rapid and requires only moderate changes in our 25 current methods (new cell-lines). Duplicate incubations are performed if kinetic studies of inhibitors need to be done in intact cells (how rapid does inhibition occur in cells). The only potential problem with this approach might be contamination of cell lysates with a compound that does not enter cells but 30 contaminates lysates during their preparation. Precautions to avoid this possible problem are taken and include studies to determine what the optimal "washing" procedure will be. The major advantage of this system is that compounds that may require labor intensive modifications (phosphorylation of nucleosides) 35 for testing with purified RDRP are rapidly screened. A more rapid screening can be achieved by transiently transfecting cells that have been incubated with potential inhibitors with a plasmid engineered to express an HCV RNA template that also encodes an

5 easily measured reporter molecule (such as secreted alkaline phosphatase or luciferase). Such a system measures HCV RDRP activity in intact cells (concentrations of inhibitions would not be diluted by lysing cells, etc. Cells in which RDRP activity is inhibited can be rapidly screened, so that large numbers of candidate inhibitors can be screened rapidly.

10 A reporter system has been devised whereby activity of r-HCV-RDRP expressed in a host cell is required for expression of a reporter gene. The host cell is transfected with a construct designed to carry the reporter coding sequence in antisense form in a structure that models the HCV replicative intermediate, when expressed as mRNA. The mRNA has, starting from the 5' end, a cap site, the reporter coding region in the 15 antisense, (-) strand, form, an HCV internal ribosome entry site (IRES) element, also in (-) strand form, a ribozyme sequence in (+) strand form, and a polyadenylation site in (+) strand form. Such an mRNA, if translated, would give rise to a nonsense protein, encoded from the (-) strand of the reporter gene. 20 However, if the complementary strand is synthesized by r-HCV-RDRP, the coding sequence of the (+) strand is translatable as the reporter protein (e.g., luciferase, fluorescent green protein, secreted alkaline phosphatase, etc.). The complement produced by RDRP lacks a capped 5' end, since the complement 25 synthesis occurs in the cytoplasm and capping occurs in the host cell nucleus. However, the presence of the HCV-IRES element allows cap-independent translation. (The IRES element will be situated 5' to the (+) strand coding sequence in the complementary strand). The function of the ribozyme motif is to 30 remove the polyA tail from the 3' end of the (-) strand, and incidentally to remove itself as well, prior to complementary strand synthesis by RDRP. A suitable ribozyme motif is provided, for example, by the R<sub>2</sub>89<sub>cc</sub> ribozyme of hepatitis delta virus. As 35 transcribed from an integrated DNA, the reporter in RNA can be diagrammed as



5 (-) strand of reporter (+) ribozyme



(+) IRES - (+) strand of reporter

15 Additions to the foregoing structure include providing a sequence of the HCV 3' untranslated region, which provides a secondary structure that can regulate or enhance r-HCV-RDRP activity. The action of r-HCV-RDRP permits expression of the reporter gene, such that a readily identifiable reaction product such as fluorescence, chemiluminescence or dye generation reaction. The 20 presence of such reaction products indirectly indicates the activity of the r-HCV-RDRP expressed in the host cell and therefore provides a means for observing the effects of a test compound on r-HCV-RDRP activity, *in vivo*. Inhibitors of *in vivo* r-HCV-RDRP activity are potential anti-viral agents against HCV.

25 While the invention has been disclosed in detail with respect to certain specific embodiments and examples, it will be understood that further embodiments, examples and modifications made according to one or more of the teachings, principles and 30 results disclosed herein, combined with knowledge in the art as applied by a person of ordinary skill therein all fall within the scope of the invention.

## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

5

(i) APPLICANT: Hagedorn, Curt H.  
Al, Reinoldus H.

10

(ii) TITLE OF INVENTION: Recombinant Hepatitis C Virus RNA  
Replicase

(iii) NUMBER OF SEQUENCES: 10

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15

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20 (F) ZIP: 80303

20

## (v) COMPUTER READABLE FORM:

25

(A) MEDIUM TYPE: Floppy disk  
(B) COMPUTER: IBM PC compatible  
(C) OPERATING SYSTEM: PC-DOS/MS-DOS  
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30

25

## (vi) CURRENT APPLICATION DATA:

30

(A) APPLICATION NUMBER: US  
(B) FILING DATE: 27-SEP-1996  
(C) CLASSIFICATION:

30

## (vii) PRIOR APPLICATION DATA:

35

(A) APPLICATION NUMBER: US 60/004383  
(B) FILING DATE: 27-SEP-1995

35

## (viii) ATTORNEY/AGENT INFORMATION:

40

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(C) REFERENCE/DOCKET NUMBER: 76-95US

40

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45

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## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 1788 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: double  
 (D) TOPOLOGY: not relevant

## 10 (ii) MOLECULE TYPE: cDNA to mRNA

## 15 (iii) HYPOTHETICAL: NO

## (ix) FEATURE:

15 (A) NAME/KEY: CDS  
 (B) LOCATION: 1..1782

## 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

20	ATG GCT AGC ATG TCT TAC TCT TGG ACA GGC GCA CTC GTC ACC CCG TGC	48
	Met Ala Ser Met Ser Tyr Ser Trp Thr Gly Ala Leu Val Thr Pro Cys	
	1 5 10 15	
25	GCC GCG GAA GAA CAG AAA CTG CCC ATC AAT GCA CTA AGC AAC TCG TTG	96
	Ala Ala Glu Glu Gln Lys Leu Pro Ile Asn Ala Leu Ser Asn Ser Leu	
	20 25 30	
30	CTA CGT CAC CAC AAT TTG GTG TAT TCC ACC ACC TCA CGC AGT GCT TGC	144
	Leu Arg His His Asn Leu Val Tyr Ser Thr Thr Ser Arg Ser Ala Cys	
	35 40 45	
35	CAA AGG CAG AAG AAA GTC ACA TTT GAC AGA CTG CAA GTT CTG GAC AGC	192
	Gln Arg Gln Lys Lys Val Thr Phe Asp Arg Leu Gln Val Leu Asp Ser	
	50 55 60	
40	CAT TAC CAG GAC GTA CTC AAG GAG GTT AAA GCA GCG GCG TCA AAA GTG	240
	His Tyr Gln Asp Val Leu Lys Glu Val Lys Ala Ala Ser Lys Val	
	65 70 75 80	
45	AAG GCT AAC TTG CTA TCC GTA GAG GAA GCT TGC AGC CTG ACG CCC CCA	288
	Lys Ala Asn Leu Leu Ser Val Glu Glu Ala Cys Ser Leu Thr Pro Pro	
	85 90 95	

His	Ser	Ala	Lys	Ser	Lys	Phe	Gly	Tyr	Gly	Ala	Lys	Asp	Val	Arg	Cys	336	
100															110		
5	CAT	GCC	AGA	AAG	GCC	GTA	ACC	CAC	ATC	AAC	TCC	GTG	TGG	AAA	GAC	CTT	384
	His	Ala	Arg	Lys	Ala	Val	Thr	His	Ile	Asn	Ser	Val	Trp	Lys	Asp	Leu	
	115														120	125	
10	CTG	GAA	GAC	AAT	GTA	ACA	CCA	ATA	GAC	ACT	ACC	ATC	ATG	GCT	AAG	AAC	432
	Leu	Glu	Asp	Asn	Val	Thr	Pro	Ile	Asp	Thr	Thr	Ile	Met	Ala	Lys	Asn	
	130														135	140	
15	GAG	GTT	TTC	TGC	GTT	CAG	CCT	GAG	AAG	GGG	GGT	CGT	AAG	CCA	GCT	CGT	480
	Glu	Val	Phe	Cys	Val	Gln	Pro	Glu	Lys	Gly	Gly	Arg	Lys	Pro	Ala	Arg	
	145														150	155	160
20	CTC	ATC	GTG	TTC	CCC	GAT	CTG	GGC	GTG	CGC	GTG	TGC	GAA	AAG	ATG	GCT	528
	Leu	Ile	Val	Phe	Pro	Asp	Leu	Gly	Val	Arg	Val	Cys	Glu	Lys	Met	Ala	
	165														170	175	
25	TTG	TAC	GAC	GTG	GTT	ACC	AAG	CTC	CCC	TTG	GCC	GTG	ATG	GGA	AGC	TCC	576
	Leu	Tyr	Asp	Val	Val	Thr	Lys	Leu	Pro	Leu	Ala	Val	Met	Gly	Ser	Ser	
	180														185	190	
30	TAC	GGA	TTC	CAA	TAC	TCA	CCA	GGA	CAG	CGG	GTT	GAA	TTC	CTC	GTG	CAA	624
	Tyr	Gly	Phe	Gln	Tyr	Ser	Pro	Gly	Gln	Arg	Val	Glu	Phe	Leu	Val	Gln	
	195														200	205	
35	GCG	TGG	AAG	TCC	AAG	AAA	ACC	CCA	ATG	GGG	TTC	TCG	TAT	GAT	ACC	CGC	672
	Ala	Trp	Lys	Ser	Lys	Lys	Thr	Pro	Met	Gly	Phe	Ser	Tyr	Asp	Thr	Arg	
	210														215	220	
40	TGC	TTT	GAC	TCC	ACA	GTC	ACT	GAG	AGC	GAC	ATC	CGT	ACG	GAG	GAG	GCA	720
	Cys	Phe	Asp	Ser	Thr	Val	Thr	Glu	Ser	Asp	Ile	Arg	Thr	Glu	Glu	Ala	
	225														230	235	240
45	ATC	TAC	CAA	TGT	TGT	GAC	CTC	GAC	CCC	CAA	GCC	CGC	GTG	GCC	ATC	AAG	768
	Ile	Tyr	Gln	Cys	Cys	Asp	Leu	Asp	Pro	Gln	Ala	Arg	Val	Ala	Ile	Lys	
	245														250	255	
	TCC	CTC	ACC	GAG	AGG	CTT	TAT	GTT	GGG	GGC	CCT	CTT	ACC	AAT	TCA	AGG	816
	Ser	Leu	Thr	Glu	Arg	Leu	Tyr	Val	Gly	Gly	Pro	Leu	Thr	Asn	Ser	Arg	
	260														265	270	

GGG GAG AAC TGC GGC TAT CGC AGG TGC CGC GCG AGC GGC GTA CTG ACA	864
Gly Glu Asn Cys Gly Tyr Arg Arg Cys Arg Ala Ser Gly Val Leu Thr	
275 280 285	
 5 ACT AGC TGC GGT AAC ACC CTC ACT TGC TAC ATC AAG GCC CGG GCA GCC	912
Thr Ser Cys Gly Asn Thr Leu Thr Cys Tyr Ile Lys Ala Arg Ala Ala	
290 295 300	
 10 TGT CGA GCC GCA GGG CTC CAG GAC TGC ACC ATG CTC GTG TGT GGC GAC	960
Cys Arg Ala Ala Gly Leu Gln Asp Cys Thr Met Leu Val Cys Gly Asp	
305 310 315 320	
 15 GAC TTA GTC GTT ATC TGT GAA AGC GCG GGG GTC CAG GAG GAC GCG GCG	1008
Asp Leu Val Val Ile Cys Glu Ser Ala Gly Val Gln Glu Asp Ala Ala	
325 330 335	
 20 AGC CTG AGA GCC TTC ACG GAG GCT ATG ACC AGG TAC TCC GCC CCC CCC	1056
Ser Leu Arg Ala Phe Thr Glu Ala Met Thr Arg Tyr Ser Ala Pro Pro	
340 345 350	
 25 GGG GAC CCC CCA CAA CCA GAA TAC GAC TTG GAG CTC ATA ACA TCA TGC	1104
Gly Asp Pro Pro Gln Pro Glu Tyr Asp Leu Glu Leu Ile Thr Ser Cys	
355 360 365	
 30 TCC TCC AAC GTG TCA GTC GCC CAC GAC GGC GCT GGA AAG AGG GTC TAC	1152
Ser Ser Asn Val Ser Val Ala His Asp Gly Ala Gly Lys Arg Val Tyr	
370 375 380	
 35 TAC CTC ACC CGT GAC CCT ACA ACC CCC CTC GCG AGA GCT GCG TGG GAG	1200
Tyr Leu Thr Arg Asp Pro Thr Thr Pro Leu Ala Arg Ala Ala Trp Glu	
385 390 395 400	
 40 ACA GCA AGA CAC ACT CCA GTC AAT TCC TGG CTA GGC AAC ATA ATC ATG	1248
Thr Ala Arg His Thr Pro Val Asn Ser Trp Leu Gly Asn Ile Ile Met	
405 410 415	
 45 TTT GCC CCC ACA CTG TGG GCG AGG ATG ATA CTG ATG ACC CAT TTC TTT	1296
Phe Ala Pro Thr Leu Trp Ala Arg Met Ile Leu Met Thr His Phe Phe	
420 425 430	
 50 AGC GTC CTT ATA GCC AGG GAC CAG CTT GAA CAG GCC CTC GAT TGC GAG	1344
Ser Val Leu Ile Ala Arg Asp Gln Leu Glu Gln Ala Leu Asp Cys Glu	
435 440 445	

ATC TAC GGG GCC TGC TAC TCC ATA GAA CCA CTT GAT CTA CCT CCA ATC Ile Tyr Gly Ala Cys Tyr Ser Ile Glu Pro Leu Asp Leu Pro Pro Ile 450 455 460	1392
5 ATT CAA AGA CTC CAT GGC CTC AGC GCA TTT TCA CTC CAC AGT TAC TCT Ile Gln Arg Leu His Gly Leu Ser Ala Phe Ser Leu His Ser Tyr Ser 465 470 475 480	1440
10 CCA GGT GAA ATT AAT AGG GTG GCC GCA TGC CTC AGA AAA CTT GGG GTA Pro Gly Glu Ile Asn Arg Val Ala Ala Cys Leu Arg Lys Leu Gly Val 485 490 495	1488
15 CCG CCC TTG CGA GCT TGG AGA CAC CGG GCC CGG AGC GTC CGC GCT AGG Pro Pro Leu Arg Ala Trp Arg His Arg Ala Arg Ser Val Arg Ala Arg 500 505 510	1536
20 CTT CTG GCC AGA GGA GGC AGG GCT GCC ATA TGT GGC AAG TAC CTC TTC Leu Leu Ala Arg Gly Gly Arg Ala Ala Ile Cys Gly Lys Tyr Leu Phe 515 520 525	1584
25 AAC TGG GCA GTA AGA ACA AAG CTC AAA CTC ACT CCA ATA GCG GCC GCT Asn Trp Ala Val Arg Thr Lys Leu Lys Leu Thr Pro Ile Ala Ala Ala 530 535 540	1632
30 GGC CAG CTG GAC TTG TCC GGC TGG TTC ACG GCT GGC TAC AGC GGG GGA Gly Gln Leu Asp Leu Ser Gly Trp Phe Thr Ala Gly Tyr Ser Gly Gly 545 550 555 560	1680
35 GAC ATT TAT CAC AGC GTG TCT CAT GCC CGG CCC CGC TGG ATC TGG TTT Asp Ile Tyr His Ser Val Ser His Ala Arg Pro Arg Trp Ile Trp Phe 565 570 575	1728
TGC CTA CTC CTG CTT GCT GCA GGG GTA GGC ATC TAC CTC CTC CCC AAC Cys Leu Leu Leu Ala Ala Gly Val Gly Ile Tyr Leu Leu Pro Asn 580 585 590	1776
40 CGA TGA GGATCC Arg *	1788

(2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 594 amino acids
- (B) TYPE: amino acid
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Ala Ser Met Ser Tyr Ser Trp Thr Gly Ala Leu Val Thr Pro Cys  
1 5 10 15

15 Ala Ala Glu Glu Gln Lys Leu Pro Ile Asn Ala Leu Ser Asn Ser Leu  
20 25 30

20 Gln Arg Gln Lys Lys Val Thr Phe Asp Arg Leu Gln Val Leu Asp Ser  
50 55 60

25 His Tyr Gln Asp Val Leu Lys Glu Val Lys Ala Ala Ala Ser Lys Val  
65 70 75 80

Lys Ala Asn Leu Leu Ser Val Glu Glu Ala Cys Ser Leu Thr Pro Pro  
85 90 95

30 His Ser Ala Lys Ser Lys Phe Gly Tyr Gly Ala Lys Asp Val Arg Cys  
                   100                 105                 110

His Ala Arg Lys Ala Val Thr His Ile Asn Ser Val Trp Lys Asp Leu  
115 120 125

35 Leu Glu Asp Asn Val Thr Pro Ile Asp Thr Thr Ile Met Ala Lys Asn  
130 135 140

Glu Val Phe Cys Val Gln Pro Glu Lys Gly Gly Arg Lys Pro Ala Arg  
145 150 155 160

Leu Ile Val Phe Pro Asp Leu Gly Val Arg Val Cys Glu Lys Met Ala  
165 170 175

Leu Tyr Asp Val Val Thr Lys Leu Pro Leu Ala Val Met Gly Ser Ser  
 180 185 190

Tyr Gly Phe Gln Tyr Ser Pro Gly Gln Arg Val Glu Phe Leu Val Gln  
 5 195 200 205

Ala Trp Lys Ser Lys Lys Thr Pro Met Gly Phe Ser Tyr Asp Thr Arg  
 210 215 220

10 Cys Phe Asp Ser Thr Val Thr Glu Ser Asp Ile Arg Thr Glu Glu Ala  
 225 230 235 240

Ile Tyr Gln Cys Cys Asp Leu Asp Pro Gln Ala Arg Val Ala Ile Lys  
 15 245 250 255

Ser Leu Thr Glu Arg Leu Tyr Val Gly Gly Pro Leu Thr Asn Ser Arg  
 260 265 270

20 Gly Glu Asn Cys Gly Tyr Arg Arg Cys Arg Ala Ser Gly Val Leu Thr  
 275 280 285

Thr Ser Cys Gly Asn Thr Leu Thr Cys Tyr Ile Lys Ala Arg Ala Ala  
 290 295 300

25 Cys Arg Ala Ala Gly Leu Gln Asp Cys Thr Met Leu Val Cys Gly Asp  
 305 310 315 320

Asp Leu Val Val Ile Cys Glu Ser Ala Gly Val Gln Glu Asp Ala Ala  
 30 325 330 335

Ser Leu Arg Ala Phe Thr Glu Ala Met Thr Arg Tyr Ser Ala Pro Pro  
 340 345 350

35 Gly Asp Pro Pro Gln Pro Glu Tyr Asp Leu Glu Leu Ile Thr Ser Cys  
 355 360 365

Ser Ser Asn Val Ser Val Ala His Asp Gly Ala Gly Lys Arg Val Tyr  
 370 375 380

40 Tyr Leu Thr Arg Asp Pro Thr Thr Pro Leu Ala Arg Ala Ala Trp Glu  
 385 390 395 400

Thr Ala Arg His Thr Pro Val Asn Ser Trp Leu Gly Asn Ile Ile Met  
 45 405 410 415

Phe Ala Pro Thr Leu Trp Ala Arg Met Ile Leu Met Thr His Phe Phe  
420 425 430

5 Ser Val Leu Ile Ala Arg Asp Gln Leu Glu Gln Ala Leu Asp Cys Glu  
435 440 445

Ile Tyr Gly Ala Cys Tyr Ser Ile Glu Pro Leu Asp Leu Pro Pro Ile  
450 455 460

10 Ile Gln Arg Leu His Gly Leu Ser Ala Phe Ser Leu His Ser Tyr Ser  
465 470 475 480

15 Pro Gly Glu Ile Asn Arg Val Ala Ala Cys Leu Arg Lys Leu Gly Val  
485 490 495

Pro Pro Leu Arg Ala Trp Arg His Arg Ala Arg Ser Val Arg Ala Arg  
500 505 510

20 Leu Leu Ala Arg Gly Arg Ala Ala Ile Cys Gly Lys Tyr Leu Phe  
515 520 525

Asn Trp Ala Val Arg Thr Lys Leu Lys Leu Thr Pro Ile Ala Ala Ala  
530 535 540

25 Gly Gln Leu Asp Leu Ser Gly Trp Phe Thr Ala Gly Tyr Ser Gly Gly  
545 550 555 560

Asp Ile Tyr His Ser Val Ser His Ala Arg Pro Arg Trp Ile Trp Phe  
565 570 575

30 Cys Leu Leu Leu Ala Ala Gly Val Gly Ile Tyr Leu Leu Pro Asn  
580 585 590

35 Arg \*

(2) INFORMATION FOR SEQ ID NO:3:

40 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 29 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(A) DESCRIPTION: /desc = "Oligonucleotide primer"

(iii) HYPOTHETICAL: NO

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

10

ATAGCTAGCA TGTCTTACTC TTGGAGAGG

29

(2) INFORMATION FOR SEQ ID NO:4:

15

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: other nucleic acid

(A) DESCRIPTION: /desc = "Oligonucleotide primer"

(iii) HYPOTHETICAL: NO

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

30

ATAGGATCCT CATCGGTTGG GGAGGGAGG

28

(2) INFORMATION FOR SEQ ID NO:5:

35

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 5 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

40

(ii) MOLECULE TYPE: protein

(iii) HYPOTHETICAL: NO

45

(v) FRAGMENT TYPE: N-terminal

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

Ala Ser Met Ser Tyr

1 5

5

## (2) INFORMATION FOR SEQ ID NO:6:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 6 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

(ii) MOLECULE TYPE: protein

15

(iii) HYPOTHETICAL: NO

(v) FRAGMENT TYPE: N-terminal

20

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

Met Ala Ser Met Ser Tyr

1 5

25

## (2) INFORMATION FOR SEQ ID NO:7:

## (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 4 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

35

(ii) MOLECULE TYPE: protein

(iii) HYPOTHETICAL: NO

40

(v) FRAGMENT TYPE: N-terminal

## (xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Ser Met Ser Tyr

45 1

## (2) INFORMATION FOR SEQ ID NO:8:

## (i) SEQUENCE CHARACTERISTICS:

5  
(A) LENGTH: 8 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: not relevant  
(D) TOPOLOGY: unknown

10  
(ii) MOLECULE TYPE: peptide

15  
(iii) HYPOTHETICAL: YES

20  
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

25  
Ala Ser Met Ser Tyr Ser Trp Thr  
1 5

## (2) INFORMATION FOR SEQ ID NO:9:

## (i) SEQUENCE CHARACTERISTICS:

30  
(A) LENGTH: 104 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: double  
(D) TOPOLOGY: not relevant

35  
(ii) MOLECULE TYPE: other nucleic acid

(iii) HYPOTHETICAL: NO

40  
(ix) FEATURE:

35  
(A) NAME/KEY: CDS  
(B) LOCATION: 63..104

45  
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

40  
GGAGACCACA ACGGTTTCCC TCTAGAAATA ATTTTGTAA ACTTTAAGAA GGAGATATAC

60

AT ATG GCT AGC ATG ACT GGT GGA CAG CAA ATG GGT CGC GGA TCC  
Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser

104

45  
1 5 10

## (2) INFORMATION FOR SEQ ID NO:10:

## (i) SEQUENCE CHARACTERISTICS:

5  
5  
(A) LENGTH: 14 amino acids  
(B) TYPE: amino acid  
(D) TOPOLOGY: linear

10  
(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser  
1 5 10

We claim:

1. An expressible coding sequence encoding a modified HCV-RDRP having the nucleotide sequence

ATG- $N_x$ - (N<sup>S</sup>) (N<sup>M</sup>) (N<sup>S</sup>) (N<sup>Y</sup>) (N<sup>S</sup>) (N<sup>W</sup>) (N<sup>T</sup>(N<sup>G</sup>) (N<sup>A</sup>) - [NS5B<sub>34-1779</sub>]  
wherein

5  $N_x$  is any nucleotide sequence encoding 0-20 amino acids,

10 N<sup>S</sup> is a codon encoding serine,  
N<sup>M</sup> is a codon encoding methionine,  
N<sup>Y</sup> is a codon encoding tyrosine,  
N<sup>W</sup> is a codon encoding tryptophan,  
N<sup>T</sup> is a codon encoding threonine,  
N<sup>G</sup> is a codon encoding glutamic acid,  
N<sup>A</sup> is a codon encoding alanine,  
any codon in parentheses being optionally deleted, and  
any of 0-5 codons in parentheses being optionally mutated  
and [NS5B<sub>34-1779</sub>] is the nucleotide sequence of NS5B from  
positions 34-1779 of SEQ ID NO:1

15

20 2. The coding sequence of claim 1 wherein 1-5 codons in parentheses is mutated to encode alanine.

25 3. The coding sequence of claim 1 wherein 1-5 codons in parentheses is mutated to encode histidine.

30 4. The coding sequence of claim 1 wherein NS5B<sub>34-1779</sub> bears a mutation encoding an amino acid replacement selected from the group Arg<sub>570</sub>-His, Arg<sub>572</sub>-His, or Try<sub>573</sub>-His, numbered according to SEQ ID NO:2

35 5. The coding sequence of claim 1 wherein  
 $N_x$  is CGA and none of the codons in parentheses is deleted.

6. The coding sequence of claim 1 wherein  $N_x$  encodes a specific proteolytic cleavage site.

7. A recombinant HCV-RDRP comprising the amino acid sequence  
MX<sub>0-20</sub> (S) (M) (S) (Y) (S) (W) (T) (G) (A) - [A<sup>9</sup>NS5B]  
wherein

5 X<sub>1-20</sub> is any amino acid sequence of 1-20 amino acids,  
any amino acid in parentheses is optionally deleted,  
and any of 0-5 amino acids in parentheses is  
optionally replaced by an amino acid, and  
A<sup>9</sup>NS5B is the amino acid sequence of amino acids 12-  
593 of SEQ ID NO:2.

10 8. Recombinant HCV-RDRP of claim 7 in purified form.

9. The recombinant HCV-RDRP of claim 7 wherein 1-5 amino  
acids in parentheses is replaced by alanine.

15 10. the recombinant HCV-RDRP of claim 7 wherein 1-5 amino  
acids in parentheses is replaced by histidine.

20 11. The recombinant HCV-RDRP of claim 7 wherein A<sup>9</sup>NS5B  
contains an amino acid replacement selected from the  
group

Arg<sub>570</sub>-His, Arg<sub>572</sub>-His or Try<sub>573</sub>-His, numbered according to  
SEQ ID NO:2.

25 12. The recombinant HCV-RDRP of Claim 7 wherein X<sub>1-20</sub> is  
alanine and none of the amino acids in parentheses is  
deleted.

30 13. The recombinant HCV-RDRP of claim 7 wherein X<sub>1-20</sub> comprises  
a specific proteolytic cleavage site.

35 14. A method of preparing soluble RDRP enzyme from a host  
cell transformed with DNA encoding HCV-RDRP comprising  
the steps of

sonicating concentrated host cells in a lysis buffer  
comprising a non-ionic detergent, a protease inhibitor

and a host-cell-wall hydrolyzing enzyme in an enzyme-compatible buffer,

freezing and thawing sonicated cells, producing a cell lysate, and

removing particles from said lysate, said particles being removable by centrifugation at 10,000-25,000 x g whereby the RDRP enzyme is retained in the supernatant fraction.

5 10 15. The method of claim 14 wherein the host cell is *E. coli*.

10 16. The method of claim 14 wherein the host cell is *E. coli*, and the host-cell-wall hydrolyzing enzyme is lysozyme.

15 17. A polyclonal or monoclonal antibody to HCV-RDRP in partially purified form.

18. Antibody of claim 17 having the capability to inhibit the enzymic activity of HCV-RDRP

20 19. A mammalian cell line capable of expressing active HCV-RDRP comprising DNA encoding r-HCV-RDRP.

25 20. The mammalian cell line of claim 19 the cell line is selected from the group U937, hepG2 or PK15.

30 21. The mammalian cell line of claim 19 wherein the r-HCV-RDRP has the amino acid sequence MA[NS5B] where [NS5B] is the amino acid sequence of amino acids 2-593 of SEQ ID NO:2.

35 22. The mammalian cell line of claim 19 further comprising a reporter gene whose translation depends on the presence of active r-HCV-RDRP in the same cell.

23. The mammalian cell line of claim 22 wherein the reporter gene expresses messenger RNA comprising the antisense of RNA encoding the reporter.

5 24. A method for testing a compound for inhibitory activity against HCV-RDRP comprising measuring RNA product in an *in vitro* RNA synthesis reaction catalyzed by r-HCV-RDRP, and

10 comparing amounts of RNA synthesized in the presence and absence of the compound, whereby comparative reduction of the amount of RNA synthesized in the presence of the compound indicates that the compound has inhibitory activity against HCV-RDRP.

15 25. The method of claim 24 further comprising comparing the activity of a control enzyme in the presence and absence of the test compound, whereby inhibition of the control enzyme by the test compound indicates absence of a specific effect on r-HCV-RDRP.

20 26. A method for testing a compound for inhibitory activity against r-HCV-RDRP *in vivo* comprising preparing a mammalian cell line capable of expressing active r-HCV-RDRP comprising DNA encoding r-HCV-RDRP,

25 transfecting said cell line with a reporter gene whose translation depends on the presence of active r-HCV-RDRP in the same cell, whereby a reporter cell line is prepared, incubating the reporter cell line in the presence or absence of a test compound, whereby inhibitor of r-HCV-RDRP by the test compound results in inhibition of translation of the reporter gene.

30 35 27. The method of claim 26 wherein the reporter gene is selected from the group: luciferase, secreted alkaline phosphatase, or fluorescent green protein.

28. The method of claim 26 wherein the reporter gene is transcribed to yield a messenger RNA encoding the antisense of the reporter, whereby synthesis of a complementary RNA thereto by r-HCV-RDRP produces a sense messenger RNA translatable to yield active reporter.

5

29. The method of claim 27 wherein the transcribed messenger RNA of the reporter further comprises an internal ribosome entry site from HCV located 3' from the reporter antisense coding region, whereby synthesis of a complementary RNA thereto by r-HCV-RDRP produces a messenger RNA having the internal ribosome entry site located 5' from the sense messenger RNA coding region thereby providing cap-independent translation of the reporter.

10

30. A method for measuring or detecting antibody to HCV-RDRP in serum of an infected patient comprising  
20 contacting the antibody with r-HCV-RDRP whereby an antigen-antibody complex is formed  
separating the antigen-antibody complex from any unbound antibody or r-HCV-RDRP and  
measuring or detecting the antigen-antibody complex.

25 31. The method of claim 31 wherein r-HCV-RDRP is localized by gel electrophoresis, the gel is contacted with serum containing antibody to HCV-RDRP, and the antigen-antibody complex is detected or measured by labeled anti-human antibody.

30

## HCV GENOME/PREDICTED PROTEINS

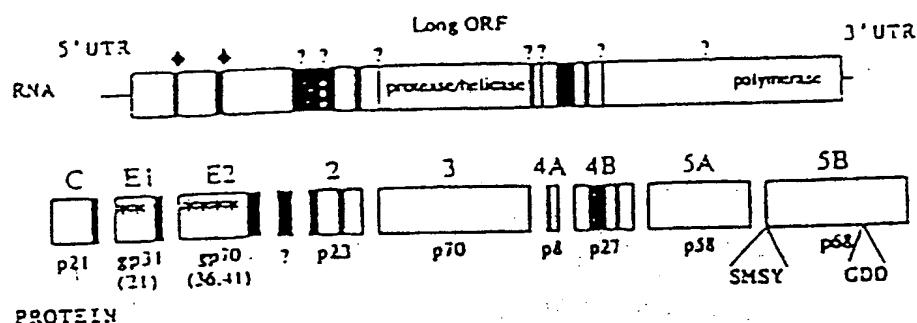


FIG. 1

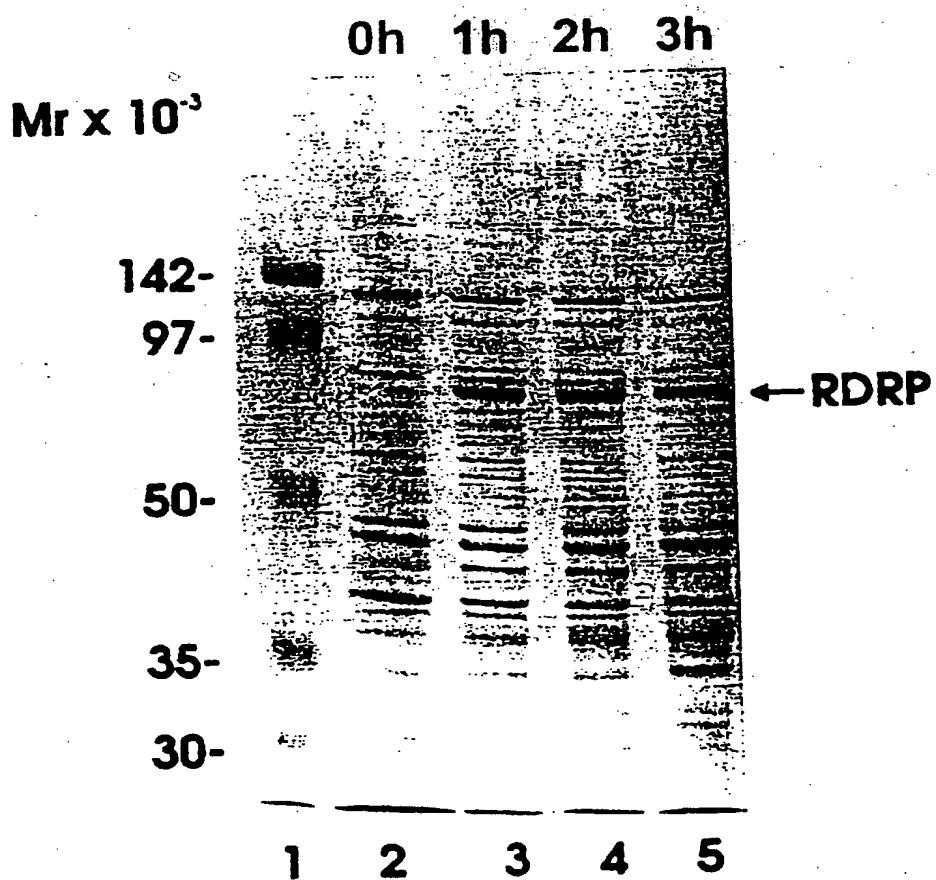


FIG. 2

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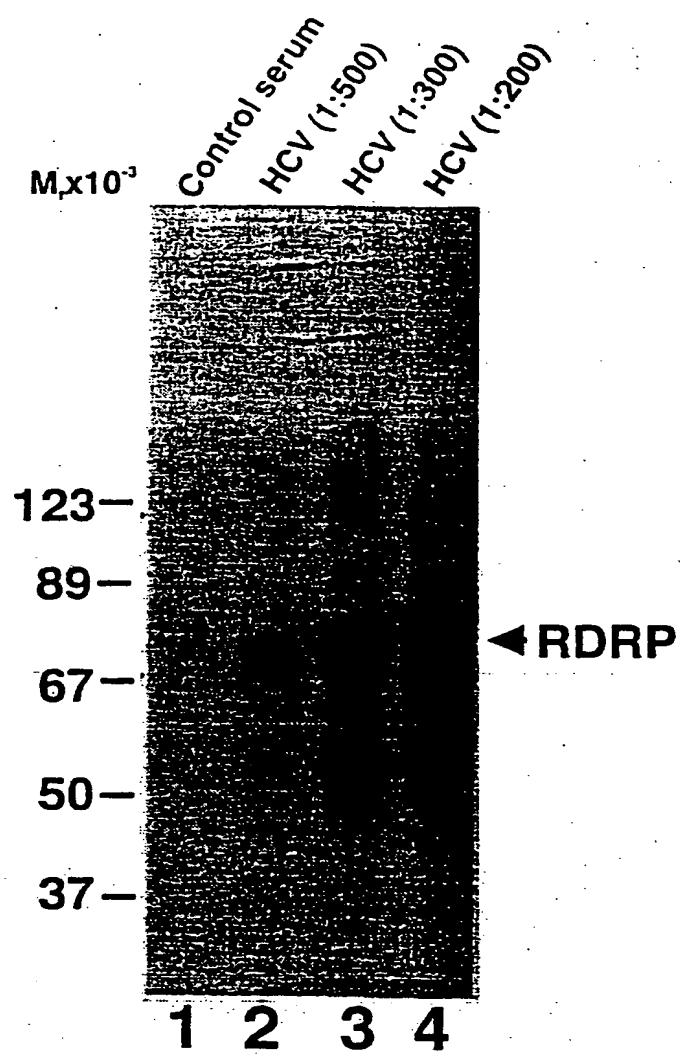


FIG. 3

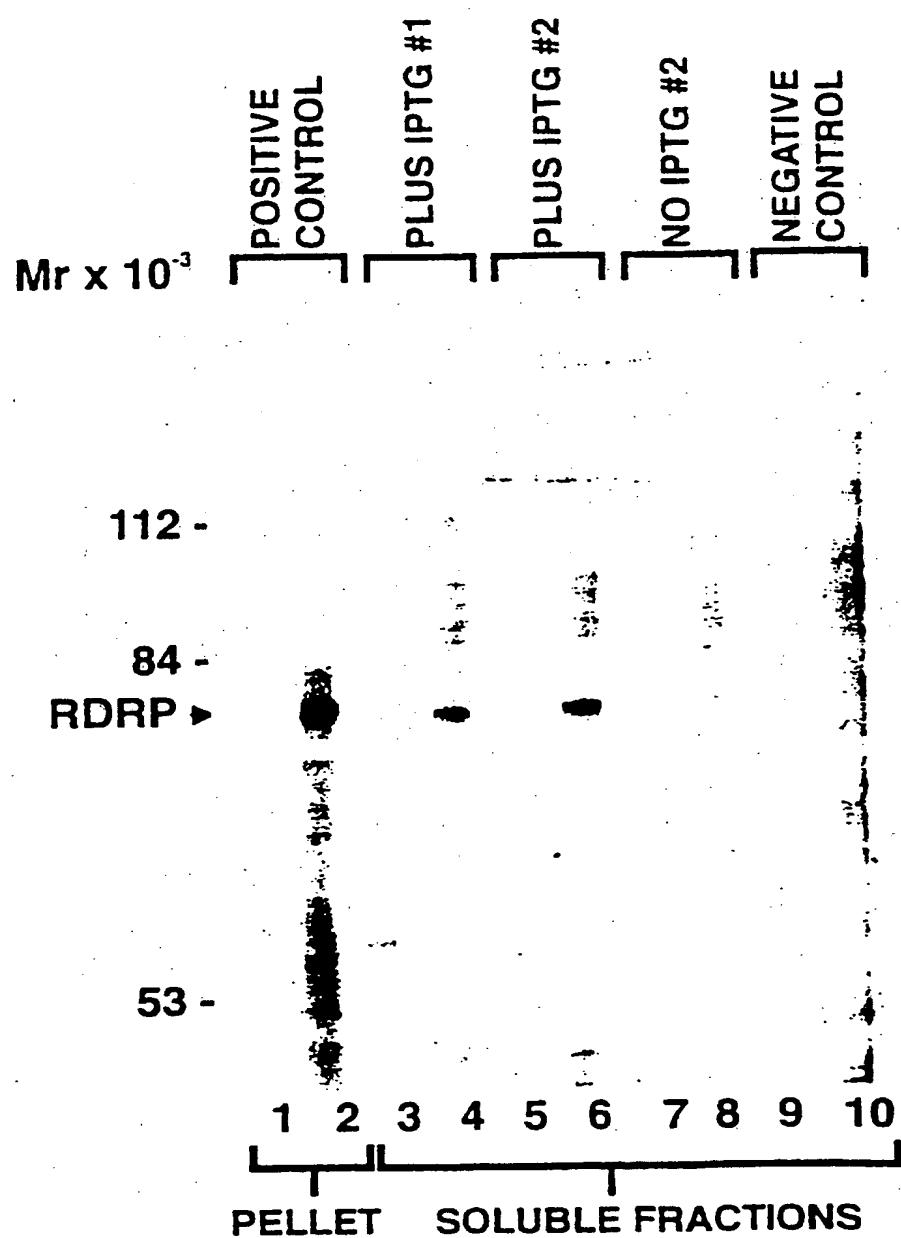


FIG. 4

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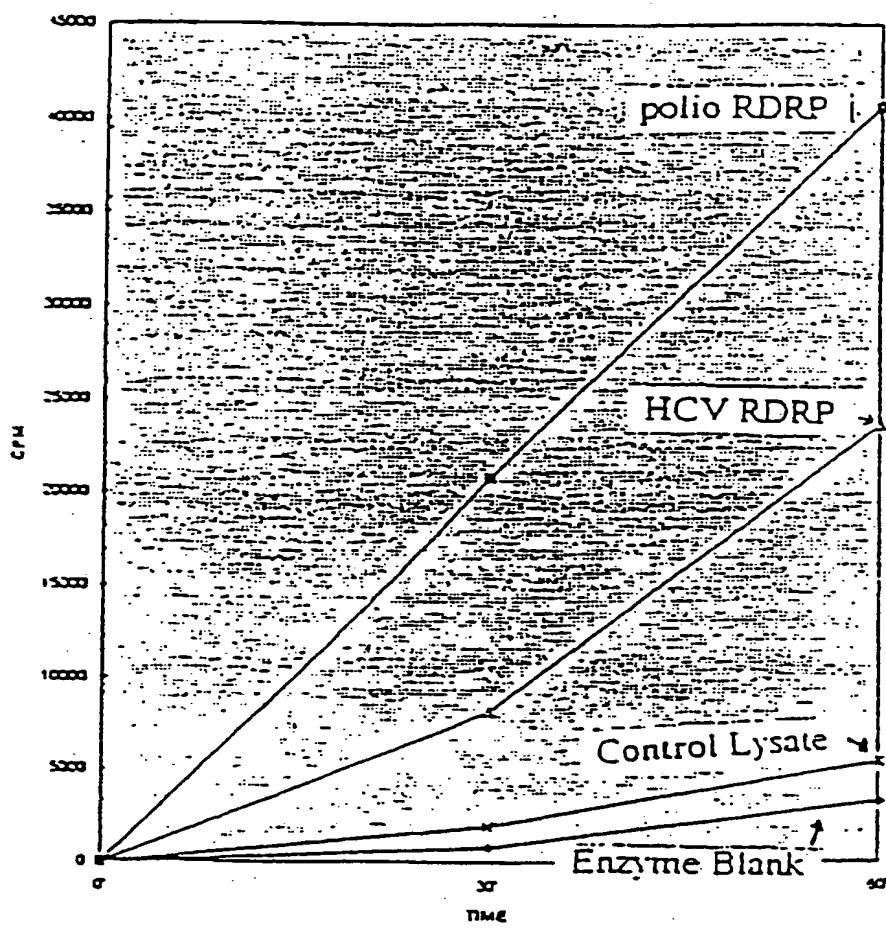
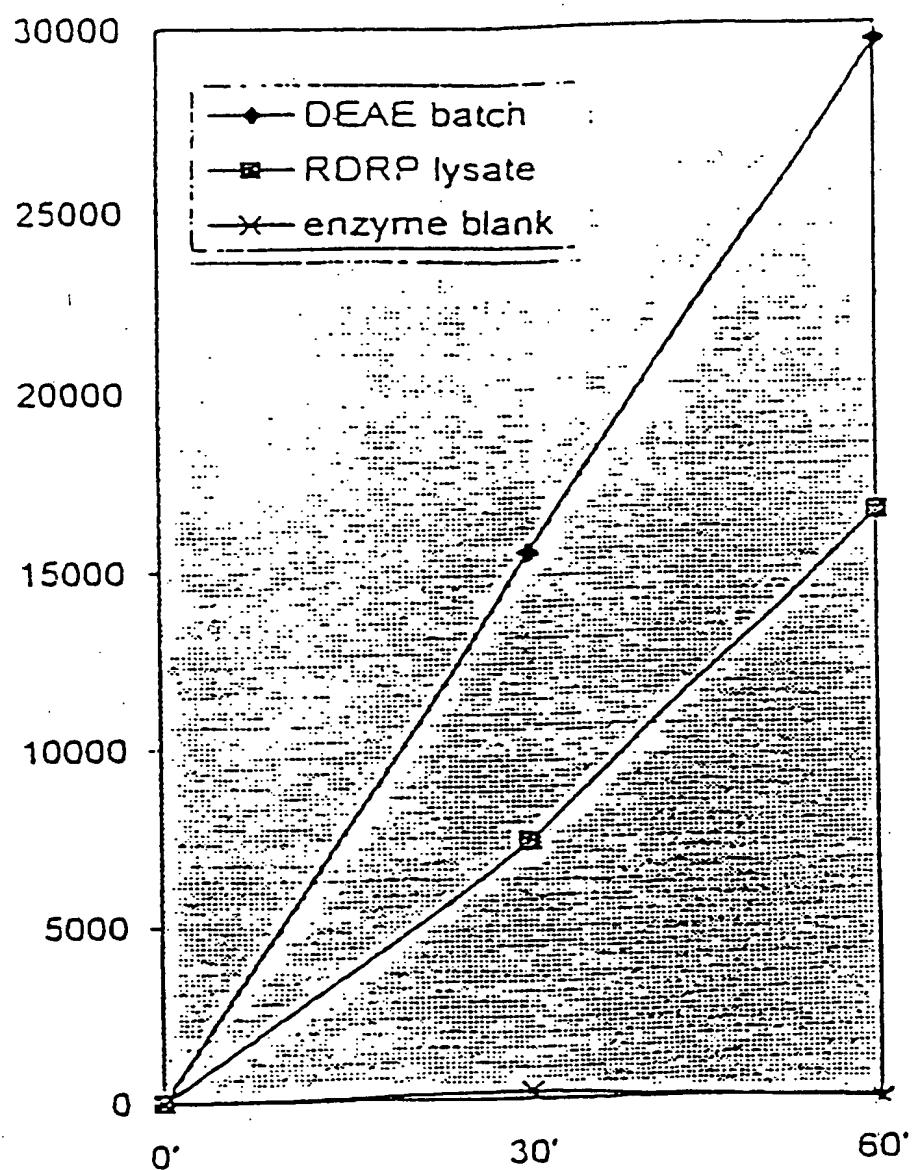


FIG. 5

**FIG. 6**

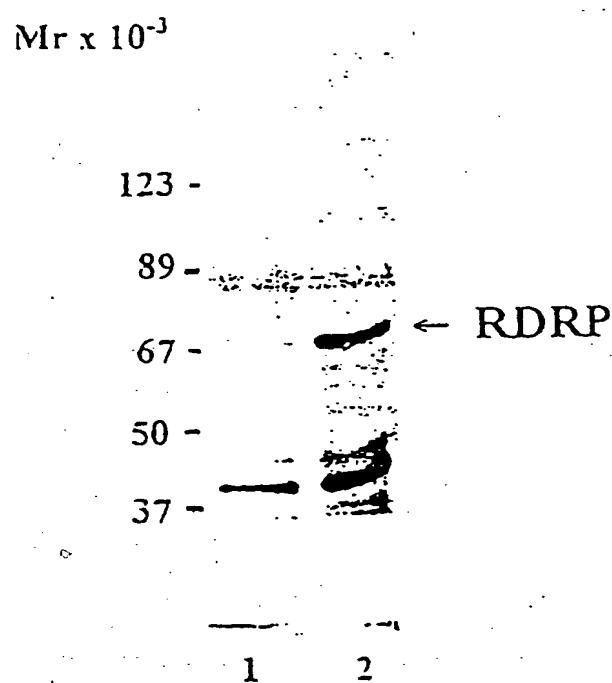


FIG. 7

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US96/15571

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C12N 9/12, 15/54; C12Q 1/48; C07K 16/00

US CL :435/194, 15, 240.2; 536/23.2, 23.4; 530/387.1

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/194, 15, 240.2; 536/23.2, 23.4; 530/387.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, DIALOG

search terms: hepatitis C virus, hcv, NS5B, RNA polymerase

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	HIROWATARI et al., Expression and processing of putative nonstructural proteins of hepatitis C virus in insect cells using baculovirus vector. Virus Research. January 1995, Vol 35, pages 43-61, see entire document.	1-31

 Further documents are listed in the continuation of Box C. See patent family annex.

•	Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
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Date of mailing of the international search report

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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>C12N 9/12, 15/54, C12Q 1/48, C07K 16/00</b>		<b>A1</b>	(11) International Publication Number: <b>WO 97/12033</b>
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(30) Priority Data: <b>60/004,383 27 September 1995 (27.09.95) US</b>			
(71) Applicant: <b>EMORY UNIVERSITY [US/US]; 2009 Ridgewood Drive, Atlanta, GA 30322 (US).</b>			
(72) Inventors: <b>HAGEDORN, Curt, H.; 4538 Club Drive, Atlanta, GA 30319 (US). AL, Reinoldus, H.; Apartment 7, 2290 LaViusta Road N.E., Atlanta, GA 30329 (US).</b>			
(74) Agents: <b>GREENLEE, Lorance, L. et al.; Greenlee, Winner and Sullivan, Suite 201, 5370 Manhattan Circle, Boulder, CO 80303 (US).</b>			

(54) Title: RECOMBINANT HEPATITIS C VIRUS RNA REPLICASE

## (57) Abstract

A recombinant RNA-dependent RNA polymerase of hepatitis C virus (r-HCV-RDRP) coding DNA was cloned and expressed yielding active enzyme *in vitro*. The r-HCV-RDRP can include up to 20 added amino acids and up to nine deleted or substituted amino acids at the NH<sub>2</sub>-terminus of the encoded amino acid sequence. The invention provides method to solubilize r-HCV-RDRP from a host cell lysate and purified r-HCV-RDRP. Methods for screening for inhibitors of r-HCV-RDRP *in vitro*, for making stably transfected mammalian cells expressing r-HCV-RDRP and for *in vivo* testing of r-HCV-RDRP inhibitors *in vivo* are disclosed. The invention provides antibodies to r-HCV-RDRP and methods for detecting antibodies to HCV-RDRP in serum of human patients.

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## RECOMBINANT HEPATITIS C VIRUS RNA REPLICASE

This application claims priority from U.S. Provisional Application Serial No. 60/004,383, filed September 27, 1995.

5

Field of the Invention

The present invention relates Hepatitis-C virus (HCV), specifically to expression and purification of an RNA-dependent RNA polymerase (RDRP) encoded by the HCV genome, to antibodies directed against HCV-RDRP and to methods of using the enzyme to diagnose chronic HCV infections and to screen for antiviral agents effective against HCV.

Background of the Invention

HCV is the major causative agent for post-transfusion and for sporadic non A, non B hepatitis (Alter, H.J. (1990) *J. Gastro. Hepatol.* 1:78-94; Dienstag, J.L. (1983) *Gastro* 85:439-462). Despite improved screening, HCV still accounts for at least 25% of the acute viral hepatitis in many countries (Alter, H.J. (1990) *supra*; Dienstag, J.L. (1983) *supra*; Alter, M.J. et al. (1990a) *J.A.M.A.* 264:2231-2235; Alter, M.J. et al (1992) *N. Engl. J. Med.* 327:1899-1905; Alter, M.J. et al .(1990b) *N. Engl. J. Med.* 321:1494-1500). Infection by HCV is insidious in a high proportion of chronically infected (and infectious) carriers who may not experience clinical symptoms for many years. The high rate of progression of acute infection to chronic infection (70-100%) and liver disease (>50%), its world-wide distribution and lack of a vaccine make HCV a significant cause of morbidity and mortality.

30

HCV is an enveloped virus whose genome is a 9.4kb single-stranded RNA (sense(+)) encoding a single polyprotein that is processed by proteolysis to yield at least 9 proteins. HCV is related to pestiviruses and flaviviruses (Choo, Q-L. et al. 5 (1989) *Science* **244**:362-364; Choo, Q-L. et al. (1991) *Proc. Natl. Acad. Sci. USA* **88**:2451-2455. Reinfestation of previously HCV-infected chimpanzees suggests that protective immunity is transient or non-existent (Farci, P. et al (1992) *Science* 10 **258**:135-140). Furthermore, results of recent vaccine trials suggest that development of an effective vaccine is remote (Houghton, M. et al. (1994) *2nd Internat. Meeting on Hepatitis C (San Diego)*). Attempted treatment of chronic HCV infection using existing antiviral agents produces low cure rates and serious side effects. (Dienstag, J.L. (1983) *supra.*)

15 The nucleotide sequence of the HCV genome has been cloned and a single open reading frame has been identified. Using a vaccinia virus expression system, several cleavage products have been tentatively identified. (Lin, C. et al. (1994) *J. Virol.* 20 **68**:5063-5073; Grakoui, A. et al. (1993) *J. Virol.* **67**:1385-1395.) The various putative cleavage products were recognized by antibodies raised against various peptides synthesized from amino acid sequences deduced from various segments of the coding regions. Sizes of antibody-reactive peptides were estimated by 25 SDS-PAGE (See Fig. 1). The non-structural protein designated 5B (NS5B) has been shown to have an amino-terminal sequence SMSY (Ser-Met-Ser-Tyr). The NS5B region encodes a 68kd protein (p68) which contains an internal GDD (Gly-Asp-Asp) motif found in RNA-dependent RNA polymerases of other RNA viruses (Koonin, E.V. 30 (1991) *J. Gen. Virol.* **72**:2197-2206). However, no polymerase activity has been detected for HCV p68. In fact, the question has been raised that the 5B protein (p68) alone does not encode 35 an active RNA-dependent RNA polymerase enzyme and that another subunit, possibly the NS5A gene product, is essential to catalytic activity. Prior attempts by the inventors and others to express the NS5B coding region as a fusion protein, using existing expression systems that facilitate purification of the

fusion product, using existing expression systems that facilitate purification of the fusion product and specific cleavage have failed to yield any active polymerase.

5 Summary of the Invention

The present invention provides a recombinant protein of HCV having RDRP activity (r-HCV-RDRP) obtainable by expression in a host mammalian or bacterial cell of a modified NS5B coding region of HCV. The modification includes addition at the amino terminus 10 of a methionine residue and optionally from 1-20 additional amino acids interposed between the N-terminal methionine and the N-terminal serine of unmodified NS5B gene product. The modification also includes deletion at the amino terminus of up to 9 amino acids to provide an amino-terminal methionine. Two 15 methionines occur naturally according to the deduced sequence of wild-type HCV-RDRP. Therefore, modification includes deletion to remove amino acids lying N-terminal to either methionine or, alternatively, deletion to some intermediate point between the two methionines plus addition of an N-terminal methionine codon. 20 A combination of deletions and insertions, within the limits described is also contemplated. Added amino acid sequence can be devised to create a specific protease cleavage site to permit post-translational modification of the recombinant HCV-RDRP expression product, *in vivo* or *in vitro*. Such post- 25 transcriptional modification can be used to generate exactly the amino acid sequence encoded by NS5B, having an N-terminal serine. Added amino acid sequence can be devised to generate an affinity ligand binding site, for convenience and ease of purification. The data reported herein were obtained with a r-HCV-RDRP having 30 an N-terminal MA (Met-Ala) dipeptide, giving an N-terminal sequence MASMSY (SEQ ID NO:6) instead of the predicted SMSY sequence of the wild-type processed protein. The coding sequence of NS5B is accordingly modified to include a met codon (ATG) at the 5'-end, as well as, optionally, codons for other amino acids 35 to be included or deleted. Minimal modifications are preferred, in order to avoid potential deleterious effects on enzyme activity, and to avoid creating artificial epitopes. The r-HCV-

RDRP can be expressed in *E. coli* and in mammalian cells to yield active RDRP. The expression of active r-HCV-RDRP in *E. coli* demonstrates that no other HCV-encoded protein is necessary for polymerase activity.

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The invention further provides r-HCV-RDRP in solubilized form, and a method of solubilization without destroying activity.

10 The invention also provides methods for purifying solubilized HCV-RDRP. One such method, to be used in combination with others, is affinity chromatography, using antibody to r-HCV-RDRP as the affinity ligand. Other affinity ligands are obtained by a combinatorial library approach as described, e.g., by Wu, J. et al. (1994) *Biochemistry* 33:14825-14833; and Ohlmeyer, 15 M.H.J. et al. (1993) *Proc. Nat. Acad. Sci. USA* 90:10922-10926.

20 In addition, the invention provides polyclonal or monoclonal antibodies specific for HCV-RDRP. Such antibodies can be made by known techniques, using the purified enzyme as antigen. Such antibodies bind either r-HCV-RDRP or wild-type HCV-RDRP. The availability of such antibodies makes it possible to prepare an affinity-labeled chromatography matrix for rapid purification of HCV-RDRP. The antibody also makes possible rapid detection of HCV-RDRP in biological materials, for example, in serum of HCV-25 infected patients.

30 The invention further provides a method for transfecting a mammalian cell with HCV-RDRP and expressing the enzyme within the cell. Consequently, the invention also provides a transfected mammalian cell line expressing r-HCV-RDRP. Such cells are useful for assaying the effects of candidate anti-viral compounds as 35 inhibitors of RDRP activity.

Therefore, the invention also provides a method for screening possible inhibitors of RDRP activity *in vivo*. Compounds with inhibitory activity can have anti-viral activity, since inhibition of the polymerase inhibits viral replication and

expression of virus gene products. The *in vitro* assay is advantageous because it can rule out compounds which cannot enter the infected cell. One class of attractive candidate compounds is the nucleoside analogs; compounds which after being modified (phosphorylated) within cells can bind to substrate sites on the enzyme or which can be incorporated into a newly synthesized RNA but whose presence there disrupts normal function of the HCV polymerase or further replication of an RNA containing the analog. Acyclovir is one example of a very effective and safe nucleoside analogue that inhibits DNA virus replication by inhibiting a viral polymerase (DNA-dependent DNA polymerase) and interfering with primer-template function (chain termination). Such analogs are almost always effective only in the nucleotide triphosphate form. The *in vitro* assay provides a convenient method of administering the compound in its nucleoside form or nucleoside monophosphate form, allowing endogenous metabolic activity of the cell to convert that form to the active triphosphate, thereby avoiding a step of chemical synthesis of the triphosphate, as would be required for an *in vivo* assay.

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A method for measuring HCV-RDRP activity *in vitro* is also provided. Such an assay permits identification of the enzyme and evaluation of its concentration during purification. In addition, the assay provides an additional, *in vitro*, method for screening potential inhibitors of RDRP as candidate anti-viral agents.

In principle, any compound can be tested as a candidate RDRP inhibitor. Certain classes of compounds are considered attractive candidates. These include, without limitation, nucleoside analogs, oligonucleotides and peptides. Certain compounds having planar, polycyclic-aromatic characteristics are also potential inhibitors. It will be understood that compounds identified as effective RDRP inhibitors must be further screened for toxicity, bioavailability, side effects and the like before being tested as therapeutic agents. Nevertheless, the initial identification as an inhibitor of HCV-RDRP is an essential first

step in the development of an anti-viral therapy. It will also be recognized that an inhibitor of r-HCV-RDRP will also inhibit wild-type HCV-RDRP.

5        In another aspect of the invention, the existence of purified HCV-RDRP or r-HCV-RDRP makes it possible to detect and measure antibodies to RDRP present in the serum of an HCV-infected patient. The fact that such antibodies exist at all is in itself a finding made possible by the expression and preparation of purified r-HCV-RDRP according to the invention. The existence of circulating antibodies to HCV-RDRP in infected serum may be due to lysis of infected cells and release of HCV-RDRP into the extracellular fluids and bloodstream, where it can stimulate an antibody response. As the disease fluctuates in severity, the amounts of HCV-RDRP released and the amounts of antibody thereto would also fluctuate. Therefore, the amount of antibody to HCV-RDRP present in a patient's serum can be used as an indicator, not only of the presence of infection, but of its severity at a given time. The assay for anti-HCV-RDRP can serve as a means of diagnosing infection and also as a means of monitoring the course of the disease over time or in response to treatment. The assay for anti-HCV-RDRP can be carried out by a variety of known techniques, such as the gel separation method described herein. Other suitable methods include ELISA, and radioimmunoassay. A sandwich-type assay, using immobilized r-HCV-RDRP to capture the antibody can then use an anti-immunoglobulin reagent tagged with an appropriate marker such as an enzyme, radioisotope, fluorescent molecule or chemiluminescent marker or the like, all as understood by those skilled in the art. (Antibodies: A laboratory manual, Ed Harlow and David Lane, Cold Spring Harbor Laboratory (1988) pp. 553-611.)

#### Brief Description of the Figures

35       Fig. 1. Hepatitis C virus genome and polyprotein cleavage products. The cleavage products of the HCV polyprotein have been tentatively identified using vaccinia virus expression systems. The amino terminus of the 5B protein expressed and processed in

5 this system is SMSY (Ser-Met-Ser-Tyr). Although published reports have not proved that the 5B protein has RNA polymerase activity, it does contain the GDD (Gly-Asp-Asp) motif found in other RNA-dependent RNA polymerases. The question has been raised that the 5B protein alone does not encode an active RNA-  
10 dependent RNA polymerase enzyme and that another subunit (possibly the NS5A gene product) is essential for catalytic activity. During the initial phases of this work we were unsure if the protein encoded by NS5B would exhibit RNA-dependent RNA polymerase activity simply due to the lack of other essential factors!

15 Fig. 2. Expression of r-HCV RNA-dependent RNA polymerase in *E. coli* using the T7 polymerase driven Studier vectors. *E. coli* containing the engineering T7 polymerase driven expression vector were incubated at 37°C until an OD<sub>600</sub> of 0.6 was reached. A sample of cells was obtained and IPTG added to a final concentration of 1 mM. Samples were collected at 1, 2 and 3 hours after IPTG induction. Whole cells were lysed in 1X sample  
20 buffer at 95°C and samples analyzed by 10% SDS-PAGE. The photograph shows a representative Coomassie Blue stained gel. Lane 1 represents molecular mass markers; lane 2, the uninduced control (0 h); lane 3, 1 h; lane 4, 2 h; and lane 5, 3 h after IPTG induction. Recombinant r-HCV RNA-dependent RNA polymerase is indicated by an arrow (RDRP).  
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30 Fig. 3. Some patients with chronic hepatitis C have circulating antibodies that react with recombinant HCV RNA-dependent RNA polymerase. Cells expressing r-HCV RDRP were harvested and lysed by heating in SDS-PAGE sample buffer. Soluble proteins were separated by SDS-PAGE, transferred to nitrocellulose membranes and immunoblotted with human sera using an Immunetics Miniblitter template (Hagedorn, et al. FEBS Lett. (1990) 264:59-62). Immunoblots were developed with a secondary 35 anti-human horseradish peroxidase conjugated antibody and enhanced chemiluminescent methods (ECL, Ambersham). This photograph shows an immunoblot where lane 1 was probed with

normal human serum and lanes 2 (1:500 dilution), 3 (1:300), and 4 (1:200) were probed with serum from a patient with chronic hepatitis C. The location of recombinant HCV RDRP (visualized by Coomassie and Ponceau S staining) is indicated by an arrow. 5 These lower molecular mass bands seen in lanes 3 & 4 represent proteolytic fragments of RDRP seen when whole *E. coli* lysates are used in immunoblots.

10 Fig. 4. Solubilization of HCV RDRP under nondenaturing conditions. Cells expressing r-HCV RDRP were harvested and processed using standard methods. Samples of insoluble *E. coli* pellets (pellet, positive control) and soluble fractions from cells containing or not containing (negative control) the RDRP expression vector were separated by SDS-PAGE and transferred to 15 nitrocellulose membranes. Proteins bound to nitrocellulose were probed with rabbit preimmune and rabbit anti-RDRP sera as outlined in Fig. 3. Blots were developed with the ECL system (Amersham). The photograph shows an immunoblot where lanes 1, 3, 5, 7 and 9 were probed with preimmune serum and lanes 2, 4, 6, 8 20 and 10 were probed with immune serum. Two independently prepared samples of soluble proteins (plus IPTC #1 & #2, lanes 3-6) were examined.

25 Fig. 5. Enzymatically active recombinant r-HCV RNA-dependent RNA polymerase. Poly(U) polymerase activity of purified recombinant poliovirus RDRP (approximately 50 ng) and equal quantities of soluble protein (approximately 1  $\mu$ g) from *E. coli* expressing r-HCV RDRP (HCV RDRP lysate) or control cells not expressing RDRP (control lysate) are shown. Incubations were 30 performed as described previously and CPM of poly(U) recovered from 15  $\mu$ l of incubation are shown at 30 min (one sample) and 60 min (mean of duplicates) of incubation are shown (J. Virol. (1986) 58:790-796).

35 Fig. 6. Enzyme assay of partially purified recombinant HCV RNA-dependent RNA polymerase. Lysates of *E. coli* expressing r-HCV RDRP were prepared and enzyme assays performed as in Fig. 5.

The experiment shown used a DEAE resin in a batch purification approach under pH and buffer conditions that allow RDRP to bind the resin. Equal quantities of protein were assayed from cell lysates (designated RDRP lysate) and proteins eluted from DEAE resin with 0.5 M NaCl (designated DEAE batch) that were concentrated to approximately that of the lysate. Additional studies with DEAE resin and other test resins have shown that partial purification of RDRP enzyme activity correlates with increases in the amount of unproteolyzed RDRP that we detect by immunoblotting using rabbit antiserum and the methods outlined in Fig. 4.

Fig. 7. Immunoaffinity purification of recombinant r-HCV RDRP. This photocopy shows an initial result with an immunoaffinity column prepared with rabbit polyclonal protein A Sepharose purified anti-RDRP antibodies. The starting material was protein solubilized from *E. coli* expressing recombinant r-HCV RDRP. The figure shows an immunoblot of proteins that were eluted from two identical columns that had protein applied under different detergent conditions. Lane 1 depicts proteins eluted from antibody/Sepharose beads that had been mixed overnight with the solubilized recombinant RDRP in 20 mM Tris-pH 7.5, 100 mM KCl, 0.5 mM EdTA, 1 mM DTT, 5% glycerol, and 0.05% Triton X-100. These beads were washed the following morning with 10 mM potassium phosphate buffer-pH 7.2 and proteins eluted with 100 mM glycine-pH 2.5. The eluted protein was collected in 1 M Tris-pH 8.0 to readjust the pH. Proteins eluted from the column were then analyzed by SDS-PAGE and immunoblotting as in Fig. 4. Lane 2 depicts proteins eluted from identical beads mixed with the same starting material except that 0.05% NP-40 was present instead of 0.05% Triton X-100. The location of r-HCV RDRP is indicated.

Detailed Description of the InventionDefinitions

"RDRP" stands for RNA-dependent RNA polymerase, an enzyme catalyzing RNA synthesis, the synthesized RNA having a sequence complementary to an RNA template. "HCV-RDRP" is the RDRP of Hepatitis C virus. The modified HCV-RDRP described herein is designated r-HCV-RDRP. The region of the HCV genome designated NS5B has been identified as a protein cleavage product of the HCV polyprotein, using a vaccinia virus expression system as described *supra*. The nucleotide sequence of NS5B is included in SEQ ID NO:1. Putative amino acid coding by the NS5B sequence begins with nucleotide 7. Where the sequence has been deleted at the 5' end, the remaining sequence has been designated by the nucleotide numbers beginning and ending the remaining coding sequences, not including the stop codon. For example NS5B<sub>34-1779</sub> designates that part of NS5B including nucleotides 34-1779 of SEQ ID NO:1.

The amino acid sequence encoding the r-HCV-RDRP exemplified herein is given in SEQ ID NO:2. The amino acid sequence encoded by NS5B begins at amino acid No. 3 of SEQ ID NO:2. Where the sequence encoded by NS5B has been deleted at the N-terminus, the remaining sequence is designated  $\Delta$ NS5B where n is the number of amino acids deleted from the N-terminus of NS5B. For example  $\Delta$ NS5B is the sequence of amino acids 12-593 in SEQ ID NO:2.

Various r-HCV-RDRP constructs are contemplated according to the invention, as described. Modified coding sequences included with the invention have the general sequence

ATG-N<sub>x</sub>- (N<sup>S</sup>) (N<sup>M</sup>) (N<sup>S</sup>) (N<sup>Y</sup>) (N<sup>S</sup>) (N<sup>W</sup>) (N<sup>T</sup>) (N<sup>G</sup>) (N<sup>A</sup>) - [NS5B<sub>34-1779</sub>] where N<sub>x</sub> is any nucleotide sequence encoding from 0-20 amino acids, N<sup>S</sup> is a codon encoding serine, N<sup>M</sup> is a codon encoding methionine, N<sup>Y</sup> is a codon encoding tyrosine, N<sup>W</sup> is a codon encoding tryptophane, N<sup>T</sup> is a codon encoding threonine, N<sup>G</sup> is a codon encoding glutamic acid, and N<sup>A</sup> is a codon encoding alanine. Any of the codons in parentheses can be deleted, if desired. Up to 5 of the codons in parentheses can be mutated if desired. The term "mutated" is

intended to mean altered to encode an amino acid other than that originally encoded by the NS5B sequence. For example, individual codons can be altered to encode alanine, by the known method of alanine scanning mutagenesis. Alanine scanning mutagenesis provides a rapid and convenient method for identifying amino acid positions where substitution is tolerated, without substantially affecting function negatively. Positions where alanine scanning reveals tolerance for substitution are likely to tolerate other amino acid substituents as well. Preferred substituents are one or more histidine residues, which can serve as affinity ligands for metal (e.g. nickel) columns. The presence of histidine provides preferential binding to the column to facilitate purification of r-HCV-RDRP. [NS5B<sub>34-1779</sub>], as defined, represents the nucleotide sequence of the remainder of SEQ ID NO:1, from nucleotides 34-1779, not including the stop codon. The techniques for making any of the foregoing sequences are essentially as described below for the sequence where N<sub>x</sub> is GCT and none of the codons in parentheses, encoding the first nine amino acids encoded by NS5B, is deleted. It will be apparent that primers can be synthesized for the desired sequence combined with desired restriction site sequences to facilitate insertion into appropriate expression vectors. The choice of vector is based on factors known in the art, including the host cell, the type of promoter desired and the presence or absence of additional sequences which could be co-expressed with the r-HCV-RDRP. The reaction condition, PCR, vector insertion and host cell growth are as described below or as well-known in the art.

Further modifications can be made to r-HCV-RDRP. A deletion of approximately 25% of the C-terminal region that did not include the Gly-Asp-Asp motif was constructed (r-HCV-RDRP-ΔC) and tested for activity. Although protein was expressed and was detectable by immunoblotting with anti-RDRP serum, lysate of *E. coli* expressing r-HCV-RDRP-ΔC had no measurable activity in either the poly(U) assay or with globin mRNA as template. However, the region of amino acids 565-572 of NS5B (565-574 of SEQ ID NO:2) near the C-terminus is considered to be exposed at

the protein surface on the basis of modeling studies. Site directed mutagenesis has been used to alter the coding to Arg 570-His, Arg 572-His and Try 573-His (using the numbering of SEQ ID NO:2). By clustering one or more additional histidine residues in a surface region near one end of the protein, purification by metal-affinity chromatography is facilitated while enzymatic activity is not substantially affected. Other predicted surface regions that can serve as sites for mutagenesis to replace an existing amino acid with histidine include amino acids 47-56, 152-159, 183-184, 210-215, 269-272, 384-391, and 43999-442 in SEQ ID NO:2.

Although the r-HCV-RDRP can be expressed in virtually any host cell type, the enzyme should preferably be in soluble form in order to be useful in *in vitro* studies, such as testing for inhibitors. When synthesized intracellularly, the enzyme is in an insoluble form in cell lysates unless steps are taken to solubilize the enzyme. In general, host cells are collected and concentrated, then lysed by means known to disrupt the host cells, for example by the use of a host-cell-wall hydrolyzing enzyme, by sonication and the like. In general a protease inhibitor is added to protect against proteolytic enzymes released by cell lysis. A non-ionic detergent can also be employed. Sonicated cells and subcellular complexes are subjected to freezing and thawing in the presence of the above-named components. Remaining particulate matter is removed by centrifugation at 10,000-35,000xg. The r-HCV-RDRP remains in the supernatant. A detailed protocol for solubilizing r-HCV-RDRP expressed in *E. coli* is described below.

Further purification of the enzyme is accomplished by techniques and expedients known in the art. These include, but are not limited to, antibody affinity chromatography, metal-binding affinity chromatography (a technique especially suited for modified forms of the enzyme having added histidine residues) as well as conventional ion-exchange columns, differential precipitation with ammonium sulfate and other methods known in

the art, not limited to the methods specifically disclosed herein. "Purified form" is used herein to mean any preparation of the enzyme having at least 4-fold greater specific activity than that measured in a solubilized cell lysate.

5

Antibodies against r-HCV-RDRP can be generated by a variety of known techniques for making monoclonal or polyclonal antibodies. Antibodies to r-HCV-RDRP also bind to HCV-RDRP of infected cells, as shown by the discovery that circulating 10 antibodies to HCV-RDRP are detectable in serum of HCV-infected patients, using r-HCV-RDRP as the antibody ligand. A variety of monoclonal antibodies can be selected, having affinity for different epitopes of r-HCV-RDRP, as known in the art. Some 15 antibodies can be inhibitory of enzyme activity. Others can have a modest affinity that facilitates binding to an antibody-affinity column and subsequent elution under conditions that do not inactivate the enzyme.

Mammalian cells are a preferred host cell for certain 20 purposes, particularly for *in vitro* screening for inhibitors of HCV-RDRP, and also for developing cell lines that can propagate HCV in cell culture. Any of the known cell lines used for transformation can, in principle, be transformed to express r-HCV-RDRP. Preferred cell lines are those of tissue origin known 25 to be infected by HCV or similar viruses, such as flaviviruses. Such cell lines include, for example, the human macrophage cell line U937, human liver-derived hepG2 cells, and the pig kidney cell line PK15. A recently-discovered segment near the 3'-end 30 of HCV-RNA has a predicted cloverleaf-type secondary structure that is a likely site of interaction for a protein or peptide (which may be host-encoded) that regulates HCV-RDRP (Tanaka, T., et al. (1995) *Biochem. Biophys. Res. Commun.* **215**:744,749). Such regulation can take the form of altering the template specificity 35 or the catalytic activity of r-HCV-RDRP as well as the wild-type enzyme. Expression of r-HCV-RDRP in cells that normally express the regulatory protein provides an *in vivo* cell system where expression of r-HCV-RDRP closely approaches the manner of

expression in HCV-infected cells. Also, the presence of active HCV-RDRP in a cell can enhance the replication of HCV introduced by infection or genomic HCV RNA introduced by transfection, by providing a "jump start" for HCV replication. Most importantly, the ability to measure r-HCV-RDRP activity in transformed cells provides an essential key to screening potential inhibitors of HCV-RDRP for their ability to inhibit the enzyme *in vivo*.

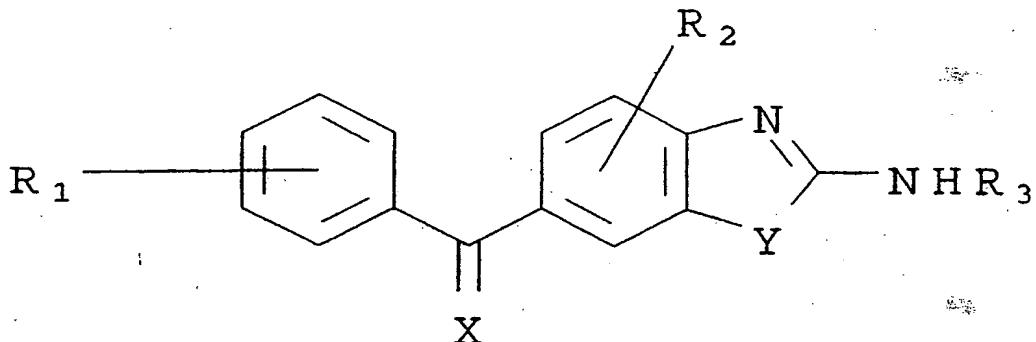
Assay for RNA synthesis by r-HCV-RDRP *in vitro* has demonstrated that the enzyme can catalyze synthesis of poly(U) using a poly(A) template, and synthesis of RNA, using a globin mRNA template. Both reactions were >90% primer-dependent under the described reaction conditions. The enzyme had no detectable activity in the absence of Mg<sup>++</sup> and showed maximal activity in the presence of about 20 mM Mg<sup>++</sup>. A method for screening compounds that have anti-viral activity is provided by testing the effect of various compounds on the RDRP enzyme activity *in vitro*. The *in vitro* method includes comparing the amounts of RNA synthesized in the presence and absence of a test compound. An inhibitory effect is indicated if the amount of RNA synthesized is reduced in a reaction where the test compound is present compared to a control reaction where the test compound is absent.

The stably transfected cell line expressing r-HCV-RDRP is especially useful for carrying out *in vivo* screening for compounds that inhibit the polymerase in intact mammalian cells. Such inhibitors are likely inhibitors of HCV replication in infected cells. An *in vivo* test is advantageous since it can screen out cytotoxic compounds, and compounds that cannot enter the cell in sufficient quantity. It also provides the added capability of testing compounds which are precursors of the actual inhibitor but converted to inhibitor by the cells metabolic processes. Examples of precursor inhibitors include nucleoside analogs which must be phosphorylated to be converted to active inhibitor, and compounds having an active group protected by a linkage that undergoes hydrolytic cleavage,

intracellularly, to form active inhibitor. Non-nucleoside analogs having the general structure shown are considered attractive candidates for r-HCV-RDRP inhibitors

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15 where R<sub>1</sub> and R<sub>2</sub> are alkyl, amino, hydroxy, alkoxy or halo groups, R<sub>3</sub> is an alkyl, aryl, hydroxy or alkoxy group, X is O, NR<sub>3</sub>, O, CH<sub>2</sub> or CHR<sub>3</sub>. In general, the test cells additionally are transformed with a reporter construct whose expression requires the action of r-HCV-RDRP, or whose expression is amplified by the presence of r-HCV-RDRP. Reporter genes are well known in the art, including, but not limited to luciferase, secreted alkaline phosphatase and the fluorescent green protein, all of which are commercially available. An attractive strategy is to use an antisense gene for the reporter, that is, a version of the reporter gene which expresses an antisense, or (-) strand messenger RNA of the reporter gene. Activity of an RDRP is then required to produce a sense (+) strand in RNA which can be translated to yield active reporter. This system has the advantage that there is no background level of reporter activity in the absence of active HCV-RDRP, if the RDRP is inhibited. An outline for the construction of suitable (-) strand reporter gene is set forth below.

## Material and Methods

35

Material - All chemicals were purchased from Fisher and all enzymes from Gibco BRL unless stated otherwise. AmpliTaq was

5 purchased from Perkin-Elmer. All other PCR and ligation components were from Invitrogen. Lysozyme, antibiotics, and pre-stained protein standards were from Sigma. Nucleotides and poly(A) were from Pharmacia. [<sup>3</sup>H-]UTP was from Dupont NEN. Oligo(U) was a generous gift from E. Ehrenfeld (University of California, Irvine).

10 Subcloning of the HCV NS5B region - PCR-primers for the amplification of the NS5B-region were designed based on the N-terminus as predicted by vaccinia virus expression studies (Lin, C. et al. (1994) *supra*; Grakoui, A. et al. (1993) *supra*) and the C-terminus based on the end of the open-reading-frame of the HCV poly protein (Choo, Q-L. et al. (1991) *supra*). The template was the original prototype HCV (type 1a) clone (obtained from the 15 CDC) (Choo, Q-L. et al. (1989), (1991), *supra*). Using the following primers, 5'-ATA GCT AGC ATG TCT TAC TCT TGG ACA GG-3' (SEQ ID NO:3) and 5'-ATA GGA TCC TCA TCG GTT GGG GAG GAG G-3' (SEQ ID NO:4), we amplified the NS5B-region with minimum changes at the N-terminus (ASMSY SEQ ID NO:5 instead of SMSY SEQ ID NO:7) and 20 directionally cloned it into pET-11a (Novagen) at *Nhe*I and *Bam*HI restriction sites (Maniatis et al. (1982) *Molecular Cloning*, Cold Spring Harbor Laboratory, Plainview, New York). The PCR amplified gene had a *Nhe*I site engineered into the 5' end and a *Bam*HI site at the 3' end. This construct results in the 25 synthesis of a recombinant protein with an amino terminal sequence of MASMSY rather than the SMSY amino terminus of the putative wild-type NS5B protein predicted by vaccinia virus expression studies.

A S M S Y S W T (amino acid encoded in primer, SEQ ID NO:8)

5'-ATA GCT AGC ATG TCT TAC TCT TGG ACA GG-B'CR primer directed at the  
 ↑  
 5 NheI cut site 5' end of the NS5B gene of the  
 hepatitis C virus cDNA,  
 SEQ ID NO:3

10

C A  
 A C  
 C = G  
 C = G  
 A - T  
 G T  
 A - T  
 G = C  
 G = C + 26  
 pppG = C

This region of the expression vector is  
 removed before inserting PCR product

15

TCTAGAAATTTTGT

IbaI

pET-11a

20

+50 +61 Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser (SEQ ID NO:10)  
 AACTTTAAGGAGATATACATATG GCT AGC ATG ACT GGT GGA CAG CAA ATG GGT CSC CGA TCC (SEQ ID NO:9)  
 SD NheI H~~he~~l BamH~~I~~

25

Restriction site in the pET-11a expression vector where the 5' end of the PCR amplified NS5B gene was inserted.

NheI

G~~X~~CTAGC

The PCR reaction was preceded by an 2 min incubation at 30 94°C, followed by 20 cycles of 1 min 94°C, 2 min 55°C, and 3 min 72°C (Coy Corporation Tempcycler II). Reactions were completed by keeping the temperature at 72°C for another 7 min and subsequent cooling to 4°C. Amplified DNA was purified by phenol/chloroform extraction, digested with NheI and BamHI and 35 re-purified using phenol/chloroform extraction and ethanol precipitation.

Insert and vector were ligated overnight at 14.5°C at an 40 3:1 ratio (insert:vector) using T4-ligase (Maniatis, *supra*). Ligated material was used to transform *E. coli* (Top10™ from Invitrogen) using CaCl<sub>2</sub> methods. Colonies were selected on ampicillin plates and minipreps of plasmid DNA isolated from single colonies were

characterized using restriction enzyme analyses. Plasmid DNA obtained by mini-preparation methods was used to transform BL21 (λDE3) *E. coli* (Novagen), organisms containing plasmid were selected using ampicillin and mini-preparations of plasmid DNA from single colonies were analyzed by restriction enzyme digestion.

10 *Expression, purification and solubilization of the putative HCV RDRP* - BL21 (λDE3) *E. coli* containing the pET-11a-NS5B construct described above were grown in overnight cultures (M9ZB media with Carbenicillin) and diluted 1:20 into fresh medium the next morning. Cells were incubated at 37°C until the culture media reached an OD<sub>600</sub> of 0.6. IPTG was added at that time to a final concentration of 1 mM. Expression of the putative RDRP was followed by SDS-PAGE analysis of whole cells lysed in sample buffer at 90°C.

20 To solubilize RDRP under non-denaturing conditions cells were harvested 2h after IPTG-induction. RDRP was solubilized by lysing the cells on ice for 20 min in 20 mM Tris pH 7.5, 100 mM KCl, 0.5 mM EDTA, 1 mM DTT, 0.1% Triton X-100 and 30 µg/ml lysozyme. Samples were sonicated on ice with an 0.5 inch probe (pulse setting) for 5 min (Ultrasonics Inc. W-225, output-setting 7) and centrifuged (19,000 g at 4°C for 30 min). The insoluble fraction (pellet) obtained from these preparations was enriched with RDRP. Pellets were suspended in SDS-PAGE sample buffer and heated for 10 min at 90°C and used as RDRP markers for SDS-PAGE gels. However, active enzyme was found in the supernatant, as set forth in the following protocol.

30

Outline of Solubilization Method for Recombinant RDRP

1. Thaw 5 g of *E. coli* pellet.

35

2. Resuspend 5 g of pellet in:

45 ml Lysis buffer  
40  $\mu$ l 100mM PMSF (plus other protease inhibitors)  
150  $\mu$ l lysozyme (10 mg/ml)

5

Lysis buffer:

20 mM Tris pH 7.5 (at 4°C)  
0.5 mM EDTA  
100 mM KCl  
1 mM DTT  
0.1% Triton X-100 (or 0.1% NP-40)  
10.0% (v/v) Glycerol

10

15

3. Place samples on ice for 20 min, then sonicate for 5 min (pulse mode; setting between 6-7). Mix while sonicating.

20

4. After sonicating flash freeze the lysate in liquid nitrogen (put the lysate into liquid nitrogen for about 1-2 min).

25

5. Quickly thaw the lysate at 37°C water bath.

30

6. Sonicate the lysate for 1 min.

35

7. Add an additional 5 ml of lysis buffer per 45 ml of sonicated sample mix.

40

8. Divide entire sonicated sample into 50 ml fractions (Fisher 50 ml tubes).

45

9. Centrifuge lysate at 12,500 rpm for 20 min in Beckman J-17 rotor (or 12,500 rpm in a Sorvall SS-34 rotor).

50

10. Remove supernatants to clean (sterile) 50 ml Fisher tubes and add sterile protein grade glycerol to a final concentration of 10% (for example, 4.44 ml of glycerol/40 ml of supernatant). This solution is stored at 4°C and used as starting material for the purification of enzymatically active HCV RDRP.

55

Further purification is accomplished by employing the following steps, either singly or in combination.

**Soluble proteins from lysate of *E. coli* expressing recombinant HCV RDRP**  
(10,000 x g supernatant)

↓

5

**45% ammonium sulfate precipitated proteins**  
(subsequently dialyzed in Tris-pH 7.5  
with 10% glycerol and 1mM DTT)

↓

10

**DEAE anion exchange chromatography**  
(starting material applied at pH 8.5 - Tris buffer,  
wash step with same buffer and enzyme  
is eluted with a 0-0.5 M NaCl gradient)

↓

15

**Phosphocellulose chromatography**

(concentrated DEAE fractions containing enzyme  
are diluted to decrease salt concentration and change  
pH to 8.0 (Tris/HCl) before applying samples to column,  
wash step, and elution with a 0-0.5 M NaCl gradient)

↓

20

**FPLC gel filtration chromatography**

(Superose 12 HR 10/30 - Pharmacia)

(starting material is concentrated, buffer changed  
to 150 mM NaCl with Tris pH 8.0 and applied in  
a 200 µl volume with a flow rate of 0.3 ml/min)

↓

25

**Recombinant HCV RDRP for enzyme assays**

(stored in aliquots at -70°C with Tris-pH 8.0,  
100 mM NaCl, 20% glycerol, 0.1% NP-40, & 1 mM DTT)

30

As in all protein purification procedures, one can modify buffers, pH and other conditions to further optimize the purification of HCV RDRP. An additional final purification step (or substitution for the FPLC gel filtration step) is a Mono-S cation exchange chromatography step at pH 6.0 with a MES buffer (the isoelectric point of the enzyme is approximately 8.8). All purification steps are monitored for enzyme activity using RDRP assay, total protein, and analyzed by SDS-PAGE.

35

Rabbit anti-HCV RDRP serum - RDRP solubilized from the pellet fraction as described above was separated by preparative SDS-PAGE and used to immunize rabbits. Animals were immunized at 4-5 week intervals as described in detail previously. (Harlow, E. and D.

Lane (1988) Antibodies: A laboratory manual, Cold Spring Harbor Laboratory, pp. 553-611.)

5 *Immunoblotting analysis* - Immunoblots were performed using previously described methods with the modification that secondary HRP conjugated antibodies were used with the enhanced chemiluminescent system (ECL, Amersham). When rabbit serum was the primary antibody, the secondary antibody was anti-rabbit immunoglobulin. When human serum was screened the secondary antibody was anti-human immunoglobulin. Serum from patients with 10 documented chronic hepatitis C infections was provided by Dr. Michael Beach of the Centers for Disease Control and Prevention (Atlanta).

15 *Poly(U) polymerase assay* - Enzyme activity in soluble fractions was measured using a poly(U) polymerase assay with poly(A) as template and oligo(U) as a primer (Hey, T.D. et al. (1986) *J. Virol.* **58**:790-796). Samples (generally 2  $\mu$ l) were assayed in 50  $\mu$ l incubations containing 50 mM HEPES (pH 8.0), 500  $\mu$ M each ATP, 20 CTP, and GTP; 4 mM DTT, 3 mM MgAc<sub>2</sub>; and 60  $\mu$ M ZnCl<sub>2</sub>. [<sup>3</sup>H]UTP at a concentration of 15  $\mu$ M (specific activity: 27 Ci/mol) was also present. Each incubation contained 1  $\mu$ g of poly(A) and 0.5  $\mu$ g oligo(U) as a primer.

25 Incubation were at 30°C for 30-60 min, [<sup>3</sup>H]poly(U) was precipitated with TCA in the presence of carrier DNA and collected on Whatman GF/C filters. Filters were washed with 0.1 M sodium pyrophosphate/1 N Hydrochloric acid and 95% ethanol, 30 respectively. [<sup>3</sup>H]poly(U) was quantitated by liquid scintillation spectrometry (LKB 1218 RackBeta).

**Stable Transfection of Baby Hamster Kidney (BHK)  
Cells Using Lipofectin**

**Day 1 (Afternoon)**

5

Split the BHK cells into 6 well plates aiming for 50% confluence for transfection

**Day 2 (after 4 p.m.)**

10

Prepare the following solutions in sterile tubes:

(A) 50  $\mu$ l miniprep DNA + 50  $\mu$ l media without serum (DMEM/F12)  
(2 each)

15

(B) 6.25  $\mu$ l Lipofectin (Life Technologies, Gaithersburg, MD) +  
93.75  $\mu$ l media

20

(C) 12.5  $\mu$ l Lipofectin + 87.5  $\mu$ l media

(D) 6.25  $\mu$ l Lipofectin + 193.75  $\mu$ l media (mock transfection)

(E) 12.5  $\mu$ l Lipofectin + 187.5  $\mu$ l media (mock transfection)

25

Gently mix A&B and A&C and let the DNA and Lipofectin react for 15 minutes at room temperature. During this time, wash the cell twice with 2 mls of DMEM/F12. Add 1.8 mls of DMEM/F12 to the DNA/Lipofectin complex and add it to the cells with gentle swirling. Leave the cells in the incubator overnight.

30

**Day 3 (9 a.m.)**

35

Remove the DNA/Lipofectin and add 3 mls of media + serum to the cells. Incubate the cells for 30-48 hours. Split the cell 1:20, 1:50 and 1:100 into 10 cm dishes in 10 mls of media + serum containing 600  $\mu$ g/ml geneticin. Allow 3-7 days for selection and 10-14 days for colony formation. The same protocol can be adapted to employ Starburst Dendrimer (Life Technologies,

Gaithersburg, MD) instead of Lipfectin, to improve transfection efficiency.

5 After selection, ring clone colonies onto 24 well plates and assay media from confluent wells for RDRP activity. Maintain cells n 600  $\mu$ g/ml geneticin.

10 Use of Stably Transfected Cells Expression HCV RDRP to Identify Compounds that Enter Intact Cells and Inhibit HCV RDRP

15 The most direct approach to determining the effect of potential inhibitors of HCV RDRP in transformed cells is to directly measure RDRP activity in cell extracts after cells have been incubated with compounds and washed extensively. This can be done using the RDRP assay described herein (with a HCV template) and requires no other new development except the cell-line. In brief, cells are incubated under conditions that maximally express active enzyme and in sufficient quantities for subsequent enzyme assays. Test compounds are added to incubations, media are removed at the desired time and cells are extensively washed to remove extracellular test compounds. Extracts of cells are prepared for RDRP assays following the general methods described herein. This approach is relatively rapid and requires only moderate changes in our current methods (new cell-lines). Duplicate incubations are performed if kinetic studies of inhibitors need to be done in intact cells (how rapid does inhibition occur in cells). The only potential problem with this approach might be contamination of cell lysates with a compound that does not enter cells but contaminates lysates during their preparation. Precautions to avoid this possible problem are taken and include studies to determine what the optimal "washing" procedure will be. The major advantage of this system is that compounds that may require labor intensive modifications (phosphorylation of nucleosides) for testing with purified RDRP are rapidly screened. A more rapid screening can be achieved by transiently transfecting cells that have been incubated with potential inhibitors with a plasmid engineered to express an HCV RNA template that also encodes an

5 easily measured reporter molecule (such as secreted alkaline phosphatase or luciferase). Such a system measures HCV RDRP activity in intact cells (concentrations of inhibitions would not be diluted by lysing cells, etc. Cells in which RDRP activity is inhibited can be rapidly screened, so that large numbers of candidate inhibitors can be screened rapidly.

10 A reporter system has been devised whereby activity of r-HCV-RDRP expressed in a host cell is required for expression of a reporter gene. The host cell is transfected with a construct designed to carry the reporter coding sequence in antisense form in a structure that models the HCV replicative intermediate, when expressed as mRNA. The mRNA has, starting from the 5' end, a cap site, the reporter coding region in the 15 antisense, (-) strand, form, an HCV internal ribosome entry site (IRES) element, also in (-) strand form, a ribozyme sequence in (+) strand form, and a polyadenylation site in (+) strand form. Such an mRNA, if translated, would give rise to a nonsense protein, encoded from the (-) strand of the reporter gene. 20 However, if the complementary strand is synthesized by r-HCV-RDRP, the coding sequence of the (+) strand is translatable as the reporter protein (e.g., luciferase, fluorescent green protein, secreted alkaline phosphatase, etc.). The complement produced by RDRP lacks a capped 5' end, since the complement 25 synthesis occurs in the cytoplasm and capping occurs in the host cell nucleus. However, the presence of the HCV-IRES element allows cap-independent translation. (The IRES element will be situated 5' to the (+) strand coding sequence in the complementary strand). The function of the ribozyme motif is to 30 remove the polyA tail from the 3' end of the (-) strand, and incidentally to remove itself as well, prior to complementary strand synthesis by RDRP. A suitable ribozyme motif is provided, for example, by the R<sub>2</sub>89<sub>cc</sub> ribozyme of hepatitis delta virus. As 35 transcribed from an integrated DNA, the reporter in RNA can be diagrammed as

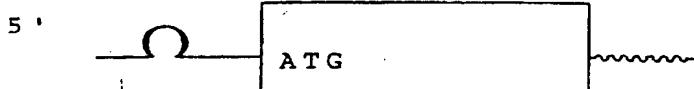
## (-) IRES



5

(-) strand of reporter (+) ribozyme

10



(+)-IRES - (+) strand of reporter

Additions to the foregoing structure include providing a sequence of the HCV 3' untranslated region, which provides a secondary structure that can regulate or enhance r-HCV-RDRP activity. The action of r-HCV-RDRP permits expression of the reporter gene, such that a readily identifiable reaction product such as fluorescence, chemiluminescence or dye generation reaction. The presence of such reaction products indirectly indicates the activity of the r-HCV-RDRP expressed in the host cell and therefore provides a means for observing the effects of a test compound on r-HCV-RDRP activity, *in vivo*. Inhibitors of *in vivo* r-HCV-RDRP activity are potential anti-viral agents against HCV.

25

While the invention has been disclosed in detail with respect to certain specific embodiments and examples, it will be understood that further embodiments, examples and modifications made according to one or more of the teachings, principles and results disclosed herein, combined with knowledge in the art as applied by a person of ordinary skill therein all fall within the scope of the invention.

## SEQUENCE LISTING

## (1) GENERAL INFORMATION:

5

(i) APPLICANT: Hagedorn, Curt H.  
Al, Reinoldus H.

10

(ii) TITLE OF INVENTION: Recombinant Hepatitis C Virus RNA  
Replicase

(iii) NUMBER OF SEQUENCES: 10

15

(iv) CORRESPONDENCE ADDRESS:

20

(A) ADDRESSEE: Greenlee, Winner and Sullivan, P.C.  
(B) STREET: 5370 Manhattan Circle, Suite 201  
(C) CITY: Boulder  
(D) STATE: Colorado  
(E) COUNTRY: US  
(F) ZIP: 80303

25

(v) COMPUTER READABLE FORM:

(A) MEDIUM TYPE: Floppy disk  
(B) COMPUTER: IBM PC compatible  
(C) OPERATING SYSTEM: PC-DOS/MS-DOS  
(D) SOFTWARE: PatentIn Release #1.0, Version #1.30

30

(vi) CURRENT APPLICATION DATA:

(A) APPLICATION NUMBER: US  
(B) FILING DATE: 27-SEP-1996  
(C) CLASSIFICATION:

35

(vii) PRIOR APPLICATION DATA:

(A) APPLICATION NUMBER: US 60/004383  
(B) FILING DATE: 27-SEP-1995

40

(viii) ATTORNEY/AGENT INFORMATION:

(A) NAME: Greenlee, Lorance L.  
(B) REGISTRATION NUMBER: 27,894  
(C) REFERENCE/DOCKET NUMBER: 76-95US

45

(ix) TELECOMMUNICATION INFORMATION:

(A) TELEPHONE: (303) 499-8080  
(B) TELEFAX: (303) 499-8089

## (2) INFORMATION FOR SEQ ID NO:1:

## (i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 1788 base pairs  
 (B) TYPE: nucleic acid  
 (C) STRANDEDNESS: double  
 (D) TOPOLOGY: not relevant

10 (ii) MOLECULE TYPE: cDNA to mRNA

15 (iii) HYPOTHETICAL: NO

## (ix) FEATURE:

15 (A) NAME/KEY: CDS  
 (B) LOCATION: 1..1782

## 20 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:

20 ATG GCT AGC ATG TCT TAC TCT TGG ACA GCA GGC CTC GTC ACC CCG TGC  
 Met Ala Ser Met Ser Tyr Ser Trp Thr Gly Ala Leu Val Thr Pro Cys  
 1 5 10 15

48

25 GCC GCG GAA GAA CAG AAA CTG CCC ATC AAT GCA CTA AGC AAC TCG TTG  
 Ala Ala Glu Glu Gln Lys Leu Pro Ile Asn Ala Leu Ser Asn Ser Leu  
 20 25 30

96

30 CTA CGT CAC CAC AAT TTG GTG TAT TCC ACC ACC TCA CGC AGT GCT TGC  
 Leu Arg His His Asn Leu Val Tyr Ser Thr Thr Ser Arg Ser Ala Cys  
 35 40 45

144

35 CAA AGG CAG AAG AAA GTC ACA TTT GAC AGA CTG CAA GTT CTG GAC AGC  
 Gln Arg Gln Lys Lys Val Thr Phe Asp Arg Leu Gln Val Leu Asp Ser  
 50 55 60

192

40 CAT TAC CAG GAC GTA CTC AAG GAG GTT AAA GCA GCG GCG TCA AAA GTG  
 His Tyr Gln Asp Val Leu Lys Glu Val Lys Ala Ala Ser Lys Val  
 65 70 75 80

240

45 AAG GCT AAC TTG CTA TCC GTA GAG GAA GCT TGC AGC CTG ACG CCC CCA  
 Lys Ala Asn Leu Leu Ser Val Glu Glu Ala Cys Ser Leu Thr Pro Pro  
 85 90 95

288

CAC TCA GCC AAA TCC AAG TTT GGT TAT GGG GCA AAA GAC GTC CGT TGC 336  
 His Ser Ala Lys Ser Lys Phe Gly Tyr Gly Ala Lys Asp Val Arg Cys  
 100 105 110

5 CAT GCC AGA AAG GCC GTA ACC CAC ATC AAC TCC GTG TGG AAA GAC CTT 384  
 His Ala Arg Lys Ala Val Thr His Ile Asn Ser Val Trp Lys Asp Leu  
 115 120 125

10 CTG GAA GAC AAT GTA ACA CCA ATA GAC ACT ACC ATC ATG GCT AAG AAC 432  
 Leu Glu Asp Asn Val Thr Pro Ile Asp Thr Thr Ile Met Ala Lys Asn  
 130 135 140

15 GAG GTT TTC TGC GTT CAG CCT GAG AAG GGG GGT CGT AAG CCA GCT CGT 480  
 Glu Val Phe Cys Val Gln Pro Glu Lys Gly Gly Arg Lys Pro Ala Arg  
 145 150 155 160

20 CTC ATC GTG TTC CCC GAT CTG GGC GTG CGC GTG TGC GAA AAG ATG GCT 528  
 Leu Ile Val Phe Pro Asp Leu Gly Val Arg Val Cys Glu Lys Met Ala  
 165 170 175

25 TTG TAC GAC GTG GTT ACC AAG CTC CCC TTG GCC GTG ATG GGA AGC TCC 576  
 Leu Tyr Asp Val Val Thr Lys Leu Pro Leu Ala Val Met Gly Ser Ser  
 180 185 190

30 TAC GGA TTC CAA TAC TCA CCA GGA CAG CGG GTT GAA TTC CTC GTG CAA 624  
 Tyr Gly Phe Gln Tyr Ser Pro Gly Gln Arg Val Glu Phe Leu Val Gln  
 195 200 205

35 GCG TGG AAG TCC AAG AAA ACC CCA ATG GGG TTC TCG TAT GAT ACC CGC 672  
 Ala Trp Lys Ser Lys Lys Thr Pro Met Gly Phe Ser Tyr Asp Thr Arg  
 210 215 220

40 TGC TTT GAC TCC ACA GTC ACT GAG AGC GAC ATC CGT ACG GAG GAG GCA 720  
 Cys Phe Asp Ser Thr Val Thr Glu Ser Asp Ile Arg Thr Glu Glu Ala  
 225 230 235 240

45 ATC TAC CAA TGT TGT GAC CTC GAC CCC CAA GCC CGC GTG GCC ATC AAG 768  
 Ile Tyr Gln Cys Cys Asp Leu Asp Pro Gln Ala Arg Val Ala Ile Lys  
 245 250 255

46 TCC CTC ACC GAG AGG CTT TAT GTT GGG GGC CCT CTT ACC AAT TCA AGG 816  
 Ser Leu Thr Glu Arg Leu Tyr Val Gly Gly Pro Leu Thr Asn Ser Arg  
 260 265 270

10	GGG GAG AAC TGC GGC TAT CGC AGG TGC CGC GCG AGC GGC GTA CTG ACA Gly Glu Asn Cys Gly Tyr Arg Arg Cys Arg Ala Ser Gly Val Leu Thr 275 280 285	864
5	ACT AGC TGC GGT AAC ACC CTC ACT TGC TAC ATC AAG GCC CGG GCA GCC Thr Ser Cys Gly Asn Thr Leu Thr Cys Tyr Ile Lys Ala Arg Ala Ala 290 295 300	912
10	TGT CGA GCC GCA GGG CTC CAG GAC TGC ACC ATG CTC GTG TGT GGC GAC Cys Arg Ala Ala Gly Leu Gln Asp Cys Thr Met Leu Val Cys Gly Asp 305 310 315 320	960
15	GAC TTA GTC GTT ATC TGT GAA AGC GCG GGG GTC CAG GAG GAC GCG GCG Asp Leu Val Val Ile Cys Glu Ser Ala Gly Val Gln Glu Asp Ala Ala 325 330 335	1008
20	AGC CTG AGA GCC TTC ACG GAG GCT ATG ACC AGG TAC TCC GCC CCC CCC Ser Leu Arg Ala Phe Thr Glu Ala Met Thr Arg Tyr Ser Ala Pro Pro 340 345 350	1056
25	GGG GAC CCC CCA CAA CCA GAA TAC GAC TTG GAG CTC ATA ACA TCA TGC Gly Asp Pro Pro Gln Pro Glu Tyr Asp Leu Glu Leu Ile Thr Ser Cys 355 360 365	1104
30	TCC TCC AAC GTG TCA GTC GCC CAC GAC GGC GCT GGA AAG AGG GTC TAC Ser Ser Asn Val Ser Val Ala His Asp Gly Ala Gly Lys Arg Val Tyr 370 375 380	1152
35	TAC CTC ACC CGT CCT ACA ACC CCC CTC GCG AGA GCT GCG TGG GAG Tyr Leu Thr Arg Asp Pro Thr Pro Leu Ala Arg Ala Ala Trp Glu 385 390 395 400	1200
40	ACA GCA AGA CAC ACT CCA GTC AAT TCC TGG CTA GGC AAC ATA ATC ATG Thr Ala Arg His Thr Pro Val Asn Ser Trp Leu Gly Asn Ile Ile Met 405 410 415	1248
45	TTT GCC CCC ACA CTG TGG GCG AGG ATG ATA CTG ATG ACC CAT TTC TTT Phe Ala Pro Thr Leu Trp Ala Arg Met Ile Leu Met Thr His Phe Phe 420 425 430	1296
	AGC GTC CTT ATA GCC AGG GAC CAG CTT GAA CAG GCC CTC GAT TGC GAG Ser Val Leu Ile Ala Arg Asp Gln Leu Glu Gln Ala Leu Asp Cys Glu 435 440 445	1344

ATC TAC GGG GCC TGC TAC TCC ATA GAA CCA CTT GAT CTA CCT CCA ATC	1392		
Ile Tyr Gly Ala Cys Tyr Ser Ile Glu Pro Leu Asp Leu Pro Pro Ile			
450	455	460	
5 ATT CAA AGA CTC CAT GGC CTC AGC GCA TTT TCA CTC CAC AGT TAC TCT	1440		
Ile Gln Arg Leu His Gly Leu Ser Ala Phe Ser Leu His Ser Tyr Ser			
465	470	475	480
10 CCA GGT GAA ATT AAT AGG GTG GCC GCA TGC CTC AGA AAA CTT GGG GTA	1488		
Pro Gly Glu Ile Asn Arg Val Ala Ala Cys Leu Arg Lys Leu Gly Val			
485	490	495	
15 CCG CCC TTG CGA GCT TGG AGA CAC CGG GCC CGG AGC GTC CGC GCT AGG	1536		
Pro Pro Leu Arg Ala Trp Arg His Arg Ala Arg Ser Val Arg Ala Arg			
500	505	510	
20 CTT CTG GCC AGA GGA GGC AGG GCT GCC ATA TGT GGC AAG TAC CTC TTC	1584		
Leu Leu Ala Arg Gly Gly Arg Ala Ala Ile Cys Gly Lys Tyr Leu Phe			
515	520	525	
25 AAC TGG GCA GTA AGA ACA AAG CTC AAA CTC ACT CCA ATA GCG GCC GCT	1632		
Asn Trp Ala Val Arg Thr Lys Leu Lys Leu Thr Pro Ile Ala Ala Ala			
530	535	540	
30 GGC CAG CTG GAC TTG TCC GGC TGG TTC ACG GCT GGC TAC AGC GGG GGA	1680		
Gly Gln Leu Asp Leu Ser Gly Trp Phe Thr Ala Gly Tyr Ser Gly Gly			
545	550	555	560
35 GAC ATT TAT CAC AGC GTG TCT CAT GCC CGG CCC CGC TGG ATC TGG TTT	1728		
Asp Ile Tyr His Ser Val Ser His Ala Arg Pro Arg Trp Ile Trp Phe			
565	570	575	
40 TGC CTA CTC CTG CTT GCT GCA GGG GTA GGC ATC TAC CTC CTC CCC AAC	1776		
Cys Leu Leu Leu Ala Ala Gly Val Gly Ile Tyr Leu Leu Pro Asn			
580	585	590	
CGA TGA GGATCC	1788		
Arg *			

## (2) INFORMATION FOR SEQ ID NO:2:

## (i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 594 amino acids  
(B) TYPE: amino acid  
(D) TOPOLOGY: linear

## (ii) MOLECULE TYPE: protein

## 10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

Met Ala Ser Met Ser Tyr Ser Trp Thr Gly Ala Leu Val Thr Pro Cys

1 5 10 15

15 Ala Ala Glu Glu Gln Lys Leu Pro Ile Asn Ala Leu Ser Asn Ser Leu

20 25 30

Leu Arg His His Asn Leu Val Tyr Ser Thr Thr Ser Arg Ser Ala Cys

35 40 45

20 Gln Arg Gln Lys Lys Val Thr Phe Asp Arg Leu Gln Val Leu Asp Ser

50 55 60

25 His Tyr Gln Asp Val Leu Lys Glu Val Lys Ala Ala Ala Ser Lys Val

65 70 75 80

55 Lys Ala Asn Leu Leu Ser Val Glu Glu Ala Cys Ser Leu Thr Pro Pro

85 90 95

30 His Ser Ala Lys Ser Lys Phe Gly Tyr Gly Ala Lys Asp Val Arg Cys

100 105 110

His Ala Arg Lys Ala Val Thr His Ile Asn Ser Val Trp Lys Asp Leu

115 120 125

35 Leu Glu Asp Asn Val Thr Pro Ile Asp Thr Thr Ile Met Ala Lys Asn

130 135 140

40 Glu Val Phe Cys Val Gln Pro Glu Lys Gly Gly Arg Lys Pro Ala Arg

145 150 155 160

Leu Ile Val Phe Pro Asp Leu Gly Val Arg Val Cys Glu Lys Met Ala

165 170 175

Leu Tyr Asp Val Val Thr Lys Leu Pro Leu Ala Val Met Gly Ser Ser  
 180 185 190

5 Tyr Gly Phe Gln Tyr Ser Pro Gly Gln Arg Val Glu Phe Leu Val Gln  
 195 200 205

Ala Trp Lys Ser Lys Lys Thr Pro Met Gly Phe Ser Tyr Asp Thr Arg  
 210 215 220

10 Cys Phe Asp Ser Thr Val Thr Glu Ser Asp Ile Arg Thr Glu Glu Ala  
 225 230 235 240

Ile Tyr Gln Cys Cys Asp Leu Asp Pro Gln Ala Arg Val Ala Ile Lys  
 245 250 255

15 Ser Leu Thr Glu Arg Leu Tyr Val Gly Gly Pro Leu Thr Asn Ser Arg  
 260 265 270

20 Gly Glu Asn Cys Gly Tyr Arg Arg Cys Arg Ala Ser Gly Val Leu Thr  
 275 280 285

Thr Ser Cys Gly Asn Thr Leu Thr Cys Tyr Ile Lys Ala Arg Ala Ala  
 290 295 300

25 Cys Arg Ala Ala Gly Leu Gln Asp Cys Thr Met Leu Val Cys Gly Asp  
 305 310 315 320

Asp Leu Val Val Ile Cys Glu Ser Ala Gly Val Gln Glu Asp Ala Ala  
 325 330 335

30 Ser Leu Arg Ala Phe Thr Glu Ala Met Thr Arg Tyr Ser Ala Pro Pro  
 340 345 350

35 Gly Asp Pro Pro Gln Pro Glu Tyr Asp Leu Glu Leu Ile Thr Ser Cys  
 355 360 365

Ser Ser Asn Val Ser Val Ala His Asp Gly Ala Gly Lys Arg Val Tyr  
 370 375 380

40 Tyr Leu Thr Arg Asp Pro Thr Pro Leu Ala Arg Ala Ala Trp Glu  
 385 390 395 400

45 Thr Ala Arg His Thr Pro Val Asn Ser Trp Leu Gly Asn Ile Ile Met  
 405 410 415

Phe Ala Pro Thr Leu Trp Ala Arg Met Ile Leu Met Thr His Phe Phe  
 420 425 430

5 Ser Val Leu Ile Ala Arg Asp Gln Leu Glu Gln Ala Leu Asp Cys Glu  
 435 440 445

Ile Tyr Gly Ala Cys Tyr Ser Ile Glu Pro Leu Asp Leu Pro Pro Ile  
 450 455 460

10 Ile Gln Arg Leu His Gly Leu Ser Ala Phe Ser Leu His Ser Tyr Ser  
 465 470 475 480

Pro Gly Glu Ile Asn Arg Val Ala Ala Cys Leu Arg Lys Leu Gly Val  
 485 490 495

15 Pro Pro Leu Arg Ala Trp Arg His Arg Ala Arg Ser Val Arg Ala Arg  
 500 505 510

Leu Leu Ala Arg Gly Arg Ala Ala Ile Cys Gly Lys Tyr Leu Phe  
 20 515 520 525

Asn Trp Ala Val Arg Thr Lys Leu Lys Leu Thr Pro Ile Ala Ala Ala  
 530 535 540

25 Gly Gln Leu Asp Leu Ser Gly Trp Phe Thr Ala Gly Tyr Ser Gly Gly  
 545 550 555 560

Asp Ile Tyr His Ser Val Ser His Ala Arg Pro Arg Trp Ile Trp Phe  
 565 570 575

30 Cys Leu Leu Leu Leu Ala Ala Gly Val Gly Ile Tyr Leu Leu Pro Asn  
 580 585 590

35 Arg \*

(2) INFORMATION FOR SEQ ID NO:3:

40 (i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 29 base pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: other nucleic acid

(A) DESCRIPTION: /desc = "Oligonucleotide primer"

(iii) HYPOTHETICAL: NO

5

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

10

ATAGCTAGCA TGTCTTACTC TTGGAGAGG

29

(2) INFORMATION FOR SEQ ID NO:4:

15

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 28 base-pairs
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS: single
- (D) TOPOLOGY: linear

20

(ii) MOLECULE TYPE: other nucleic acid

(A) DESCRIPTION: /desc = "Oligonucleotide primer"

(iii) HYPOTHETICAL: NO

25

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

30

ATAGGATCCT CATCGGTTGG GGAGGGAGG

28

(2) INFORMATION FOR SEQ ID NO:5:

35

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 5 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

40

(ii) MOLECULE TYPE: protein

(iii) HYPOTHETICAL: NO

45

(v) FRAGMENT TYPE: N-terminal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

Ala Ser Met Ser Tyr  
1 5

5

(2) INFORMATION FOR SEQ ID NO:6:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 6 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

(ii) MOLECULE TYPE: protein

15

(iii) HYPOTHETICAL: NO

(v) FRAGMENT TYPE: N-terminal

20

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

25

Met Ala Ser Met Ser Tyr  
1 5

(2) INFORMATION FOR SEQ ID NO:7:

30

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 4 amino acids
- (B) TYPE: amino acid
- (C) STRANDEDNESS: not relevant
- (D) TOPOLOGY: unknown

35

(ii) MOLECULE TYPE: protein

(iii) HYPOTHETICAL: NO

40

(v) FRAGMENT TYPE: N-terminal

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

Ser Met Ser Tyr  
1

45

## (2) INFORMATION FOR SEQ ID NO:8:

## (i) SEQUENCE CHARACTERISTICS:

5 (A) LENGTH: 8 amino acids  
(B) TYPE: amino acid  
(C) STRANDEDNESS: not relevant  
(D) TOPOLOGY: unknown

## 10 (ii) MOLECULE TYPE: peptide

## (iii) HYPOTHETICAL: YES

## 15 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

Ala Ser Met Ser Tyr Ser Trp Thr  
1 5

## 20 (2) INFORMATION FOR SEQ ID NO:9:

## (i) SEQUENCE CHARACTERISTICS:

25 (A) LENGTH: 104 base pairs  
(B) TYPE: nucleic acid  
(C) STRANDEDNESS: double  
(D) TOPOLOGY: not relevant

## 30 (ii) MOLECULE TYPE: other nucleic acid

## (iii) HYPOTHETICAL: NO

## 35 (ix) FEATURE:

(A) NAME/KEY: CDS  
(B) LOCATION: 63..104

## 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

45 GGAGACCACA ACGGTTTCCC TCTAGAAATA ATTTTGTAA ACTTTAAGAA GGAGATATAC 60

AT ATG GCT AGC ATG ACT GGT GGA CAG CAA ATG GGT CGC GGA TCC  
Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser

45 1 5 10

## (2) INFORMATION FOR SEQ ID NO:10:

## (i) SEQUENCE CHARACTERISTICS:

5  
(A) LENGTH: 14 amino acids  
(B) TYPE: amino acid  
(D) TOPOLOGY: linear

10  
(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:

Met Ala Ser Met Thr Gly Gly Gln Gln Met Gly Arg Gly Ser  
1 5 10

We claim:

1. An expressible coding sequence encoding a modified HCV-RDRP having the nucleotide sequence

ATG-N<sub>x</sub>- (N<sup>S</sup>) (N<sup>M</sup>) (N<sup>S</sup>) (N<sup>Y</sup>) (N<sup>S</sup>) (N<sup>W</sup>) (N<sup>T</sup>(N<sup>G</sup>) (N<sup>A</sup>) - [NS5B<sub>34-1779</sub>]

wherein

5 N<sub>x</sub> is any nucleotide sequence encoding 0-20 amino acids,

N<sup>S</sup> is a codon encoding serine,

N<sup>M</sup> is a codon encoding methionine,

N<sup>Y</sup> is a codon encoding tyrosine,

10 N<sup>W</sup> is a codon encoding tryptophan,

N<sup>T</sup> is a codon encoding threonine,

N<sup>G</sup> is a codon encoding glutamic acid,

N<sup>A</sup> is a codon encoding alanine,

any codon in parentheses being optionally deleted, and

15 any of 0-5 codons in parentheses being optionally mutated

and [NS5B<sub>34-1779</sub>] is the nucleotide sequence of NS5B from positions 34-1779 of SEQ ID NO:1

2. The coding sequence of claim 1 wherein 1-5 codons in parentheses is mutated to encode alanine.

20 3. The coding sequence of claim 1 wherein 1-5 codons in parentheses is mutated to encode histidine.

25 4. The coding sequence of claim 1 wherein NS5B<sub>34-1779</sub> bears a mutation encoding an amino acid replacement selected from the group Arg<sub>570</sub>-His, Arg<sub>572</sub>-His, or Try<sub>573</sub>-His, numbered according to SEQ ID NO:2

30 5. The coding sequence of claim 1 wherein N<sub>x</sub> is CGA and none of the codons in parentheses is deleted.

35 6. The coding sequence of claim 1 wherein N<sub>x</sub> encodes a specific proteolytic cleavage site.

7. A recombinant HCV-RDRP comprising the amino acid sequence  
 $MX_{0-20}$  (S) (M) (S) (Y) (S) (W) (T) (G) (A) - [ $\Delta^9$ NS5B]  
wherein

5  $X_{1-20}$  is any amino acid sequence of 1-20 amino acids,  
any amino acid in parentheses is optionally deleted,  
and any of 0-5 amino acids in parentheses is  
optionally replaced by an amino acid, and  
 $\Delta^9$ NS5B is the amino acid sequence of amino acids 12-  
593 of SEQ ID NO:2.

10

8. Recombinant HCV-RDRP of claim 7 in purified form.

9. The recombinant HCV-RDRP of claim 7 wherein 1-5 amino acids in parentheses is replaced by alanine.

15

10. the recombinant HCV-RDRP of claim 7 wherein 1-5 amino acids in parentheses is replaced by histidine.

20

11. The recombinant HCV-RDRP of claim 7 wherein  $\Delta^9$ NS5B contains an amino acid replacement selected from the group Arg<sub>570</sub>-His, Arg<sub>572</sub>-His or Try<sub>573</sub>-His, numbered according to SEQ ID NO:2.

25

12. The recombinant HCV-RDRP of Claim 7 wherein  $X_{1-20}$  is alanine and none of the amino acids in parentheses is deleted.

30

13. The recombinant HCV-RDRP of claim 7 wherein  $X_{1-20}$  comprises a specific proteolytic cleavage site.

35

14. A method of preparing soluble RDRP enzyme from a host cell transformed with DNA encoding HCV-RDRP comprising the steps of sonicating concentrated host cells in a lysis buffer comprising a non-ionic detergent, a protease inhibitor

and a host-cell-wall hydrolyzing enzyme in an enzyme-compatible buffer;

freezing and thawing sonicated cells, producing a cell lysate, and

removing particles from said lysate, said particles being removable by centrifugation at 10,000-25,000 x g whereby the RDRP enzyme is retained in the supernatant fraction.

5 10 15. The method of claim 14 wherein the host cell is *E. coli*.

16. The method of claim 14 wherein the host cell is *E. coli*, and the host-cell-wall hydrolyzing enzyme is lysozyme.

15 17. A polyclonal or monoclonal antibody to HCV-RDRP in partially purified form.

18. Antibody of claim 17 having the capability to inhibit the enzymic activity of HCV-RDRP

20 19. A mammalian cell line capable of expressing active HCV-RDRP comprising DNA encoding r-HCV-RDRP.

25 20. The mammalian cell line of claim 19 the cell line is selected from the group U937, hepG2 or PK15.

30 21. The mammalian cell line of claim 19 wherein the r-HCV-RDRP has the amino acid sequence MA[NS5B] where [NS5B] is the amino acid sequence of amino acids 2-593 of SEQ ID NO:2,

22. The mammalian cell line of claim 19 further comprising a reporter gene whose translation depends on the presence of active r-HCV-RDRP in the same cell.

23. The mammalian cell line of claim 22 wherein the reporter gene expresses messenger RNA comprising the antisense of RNA encoding the reporter.

5 24. A method for testing a compound for inhibitory activity against HCV-RDRP comprising measuring RNA product in an *in vitro* RNA synthesis reaction catalyzed by r-HCV-RDRP, and

10 comparing amounts of RNA synthesized in the presence and absence of the compound, whereby comparative reduction of the amount of RNA synthesized in the presence of the compound indicates that the compound has inhibitory activity against HCV-RDRP.

15 25. The method of claim 24 further comprising comparing the activity of a control enzyme in the presence and absence of the test compound, whereby inhibition of the control enzyme by the test compound indicates absence of a specific effect on r-HCV-RDRP.

20 26. A method for testing a compound for inhibitory activity against r-HCV-RDRP *in vivo* comprising preparing a mammalian cell line capable of expressing active r-HCV-RDRP comprising DNA encoding r-HCV-RDRP,

25 transfecting said cell line with a reporter gene whose translation depends on the presence of active r-HCV-RDRP in the same cell, whereby a reporter cell line is prepared, incubating the reporter cell line in the presence or absence of a test compound, whereby inhibitor of r-HCV-RDRP by the test compound results in inhibition of translation of the reporter gene.

30 27. The method of claim 26 wherein the reporter gene is selected from the group: luciferase, secreted alkaline phosphatase, or fluorescent green protein.

28. The method of claim 26 wherein the reporter gene is transcribed to yield a messenger RNA encoding the antisense of the reporter, whereby synthesis of a complementary RNA thereto by r-HCV-RDRP produces a sense messenger RNA translatable to yield active reporter.

5

29. The method of claim 27 wherein the transcribed messenger RNA of the reporter further comprises an internal ribosome entry site from HCV located 3' from the reporter antisense coding region, whereby synthesis of a complementary RNA thereto by r-HCV-RDRP produces a messenger RNA having the internal ribosome entry site located 5' from the sense messenger RNA coding region thereby providing cap-independent translation of the reporter.

10

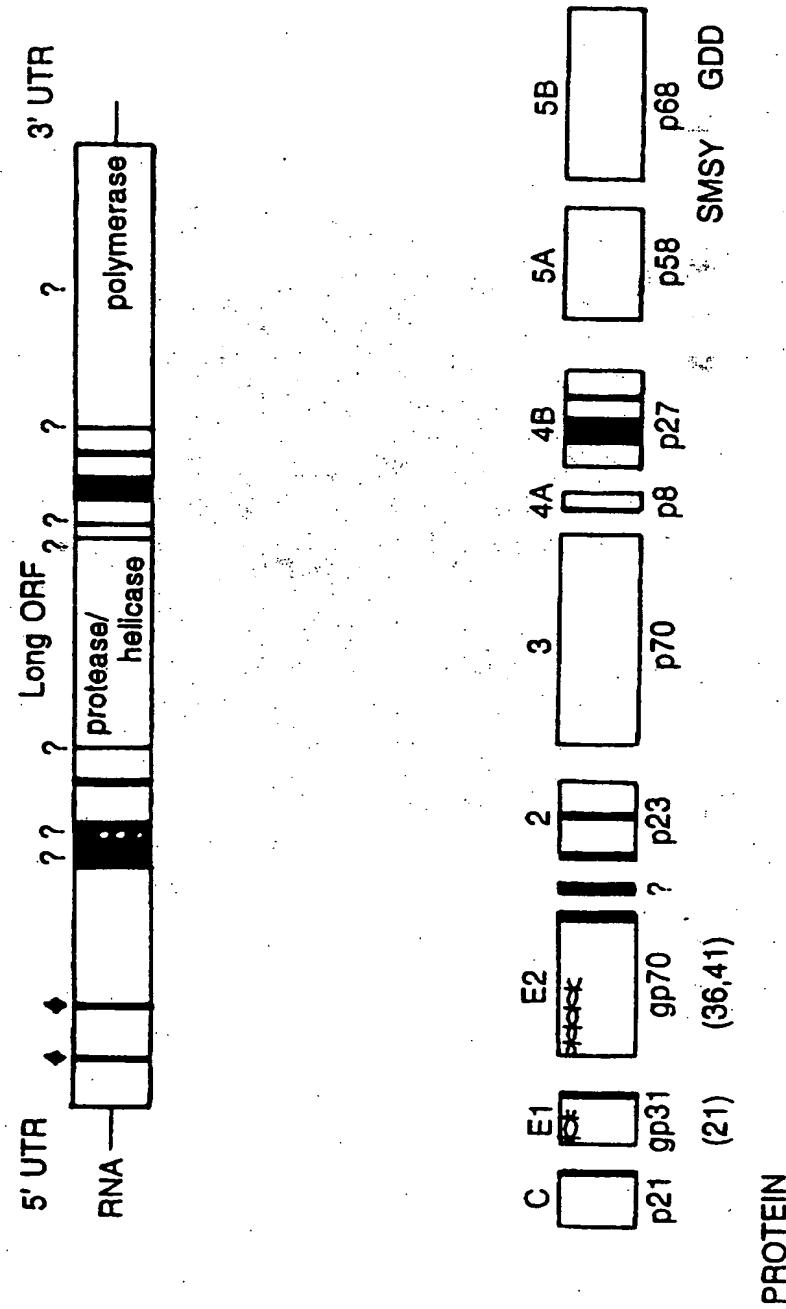
30. A method for measuring or detecting antibody to HCV-RDRP in serum of an infected patient comprising  
contacting the antibody with r-HCV-RDRP whereby an antigen-antibody complex is formed  
separating the antigen-antibody complex from any unbound antibody or r-HCV-RDRP and  
measuring or detecting the antigen-antibody complex.

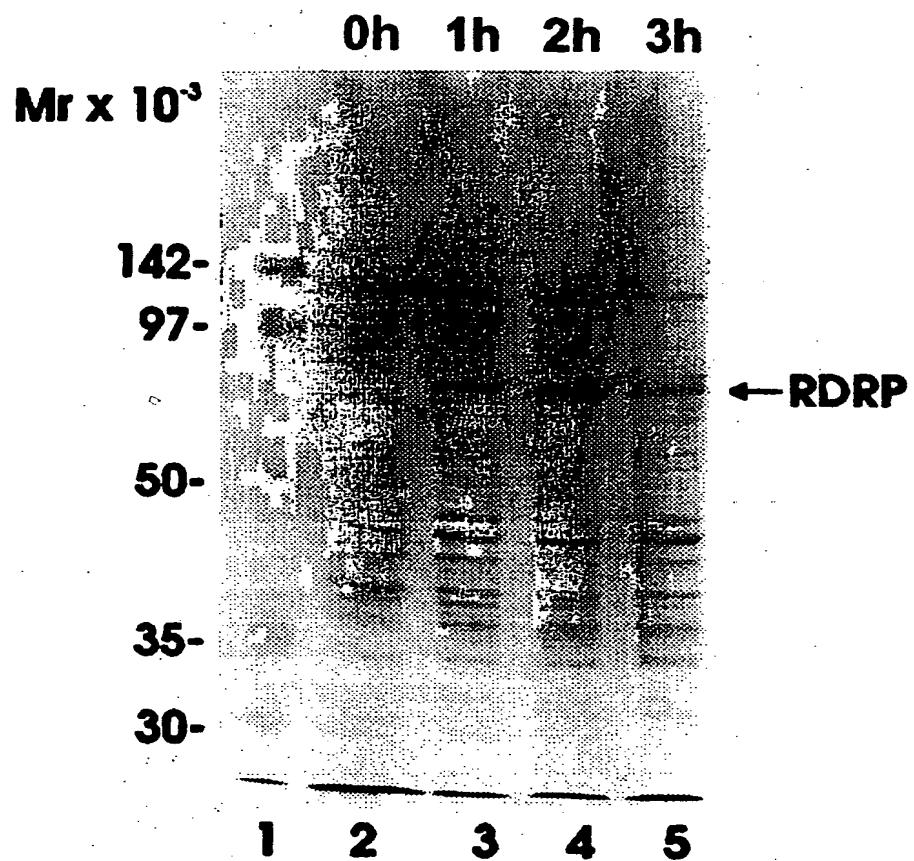
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20

25. 31. The method of claim 31 wherein r-HCV-RDRP is localized by gel electrophoresis, the gel is contacted with serum containing antibody to HCV-RDRP, and the antigen-antibody complex is detected or measured by labeled anti-human antibody.

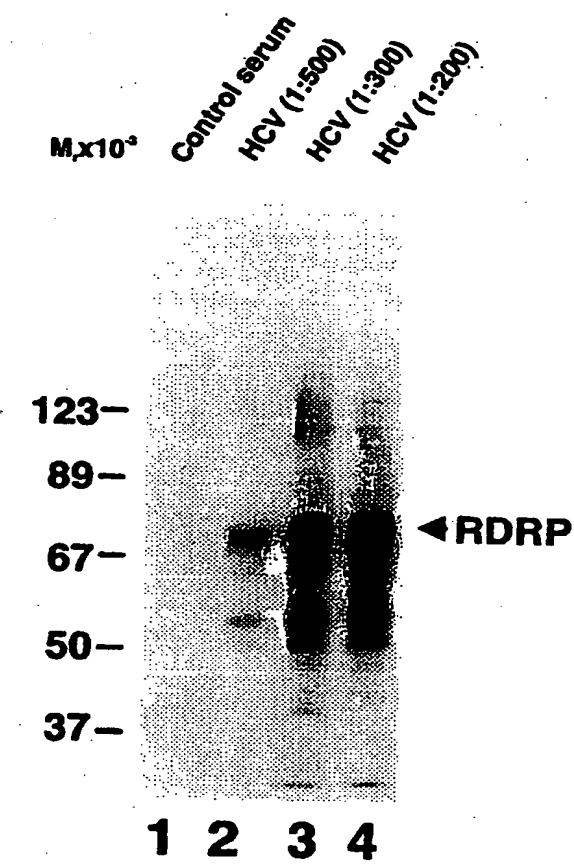
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**FIG. 1****SUBSTITUTE SHEET (RULE 26)**



**FIG. 2**

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**FIG. 3**

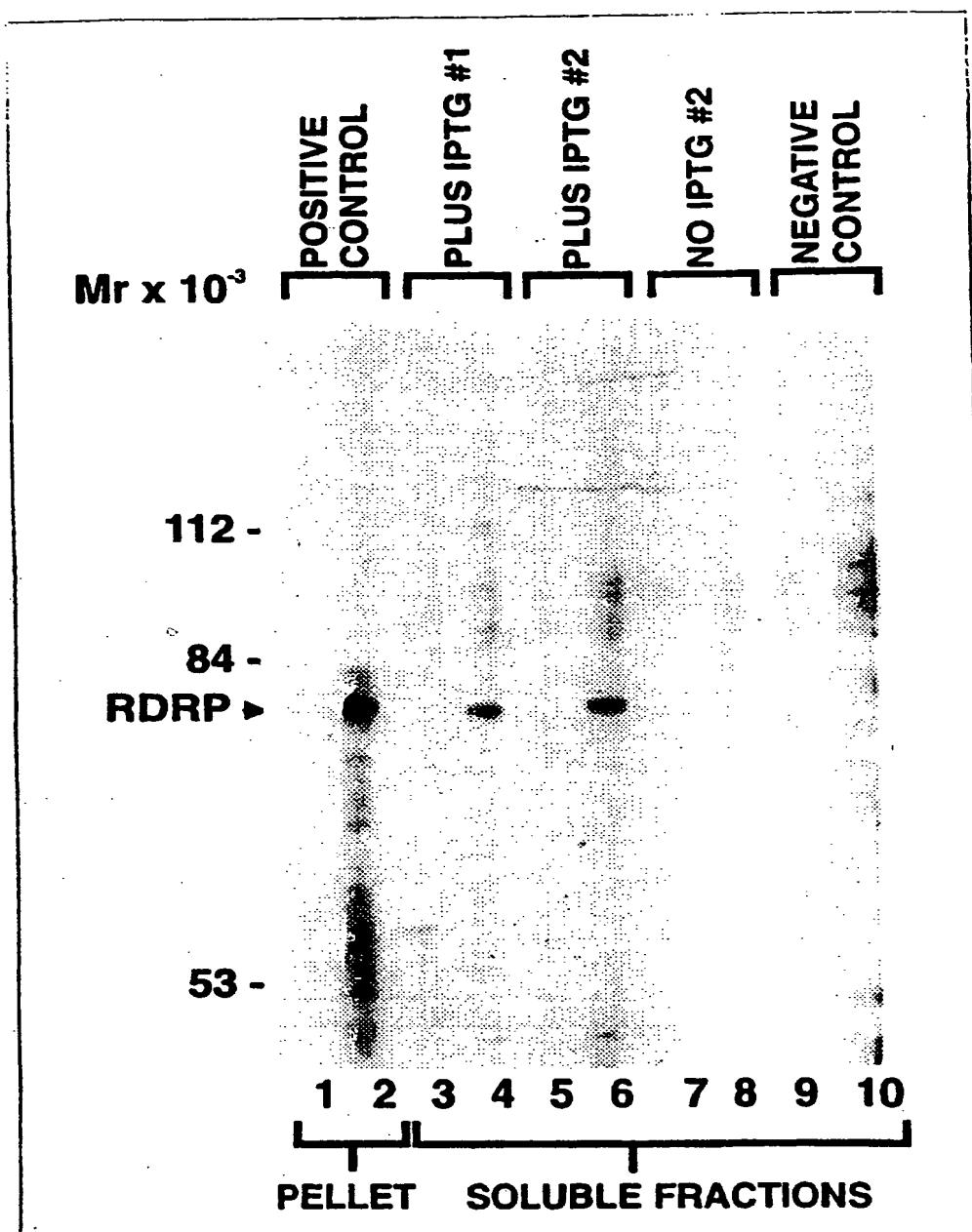
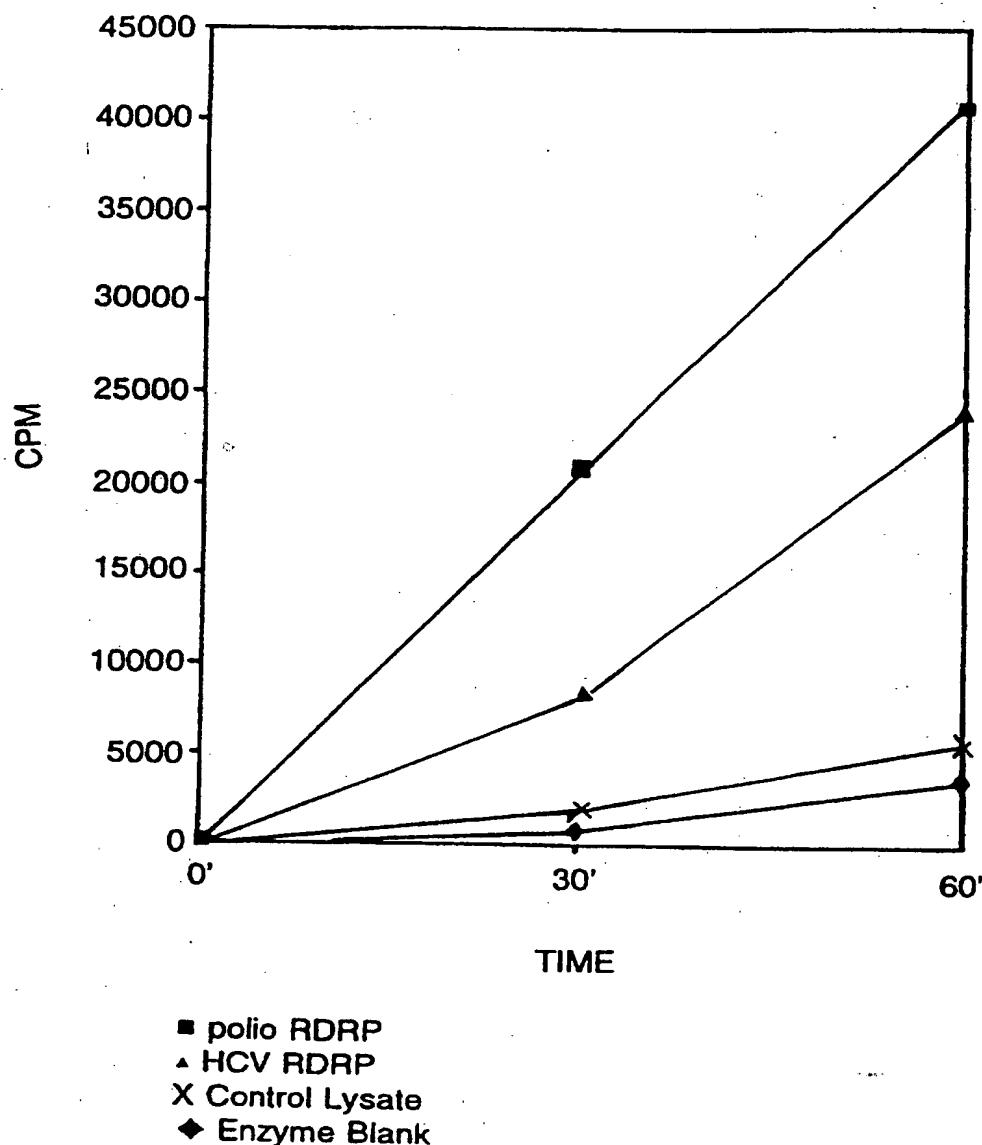
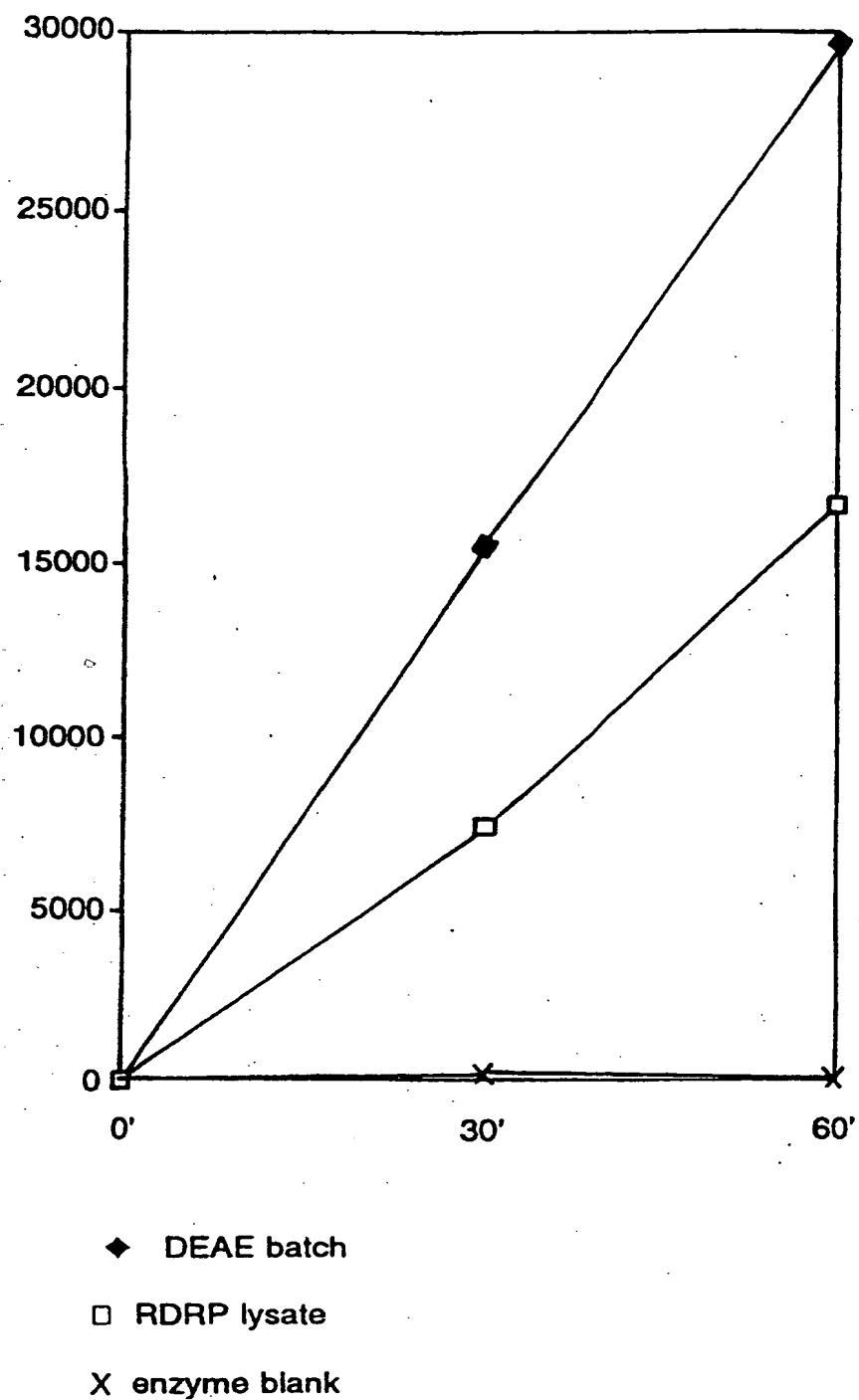
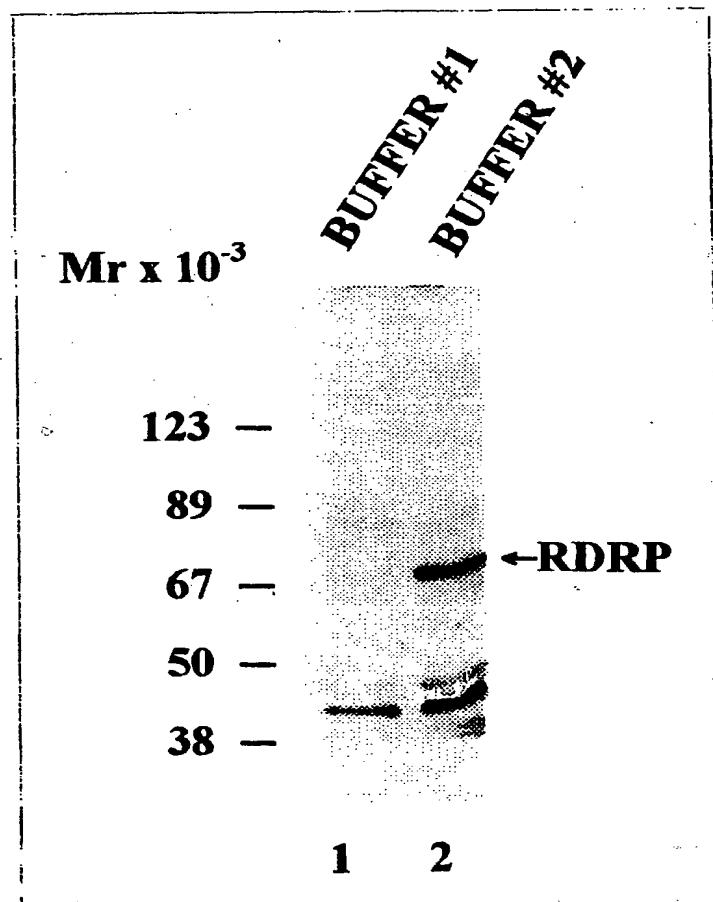


FIG. 4

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**FIG. 5****SUBSTITUTE SHEET (RULE 26)**

**FIG. 6****SUBSTITUTE SHEET (RULE 26)**

**FIG. 7**

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/15571

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C12N 9/12, 15/54; C12Q 1/48; C07K 16/00  
US CL :435/194, 15, 240.2; 536/23.2, 23.4; 530/387.1

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/194, 15, 240.2; 536/23.2, 23.4; 530/387.1

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, DIALOG  
search terms: hepatitis C virus, hcv, NS5B, RNA polymerase

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	HIROWATARI et al., Expression and processing of putative nonstructural proteins of hepatitis C virus in insect cells using baculovirus vector. Virus Research. January 1995, Vol 35, pages 43-61, see entire document.	1-31

Further documents are listed in the continuation of Box C.  See patent family annex.

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Date of the actual completion of the international search

20 NOVEMBER 1996

Date of mailing of the international search report

26 DEC 1996

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